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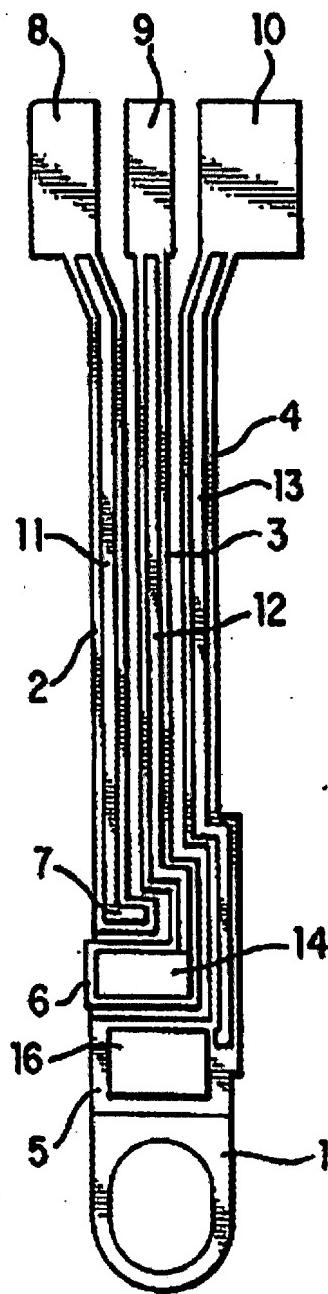
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The present invention is based on the discovery of NAD<+> and NADP<+> mediator compounds that do not bind irreversibly to thiol groups in the active sites of intracellular dehydrogenase enzymes. Such mediator compounds avoid a common mode of enzyme inhibition. The mediators can therefore increase the stability and reliability of the electrical response in amperometric electrodes constructed from NAD- or NADP-dependent enzymes.



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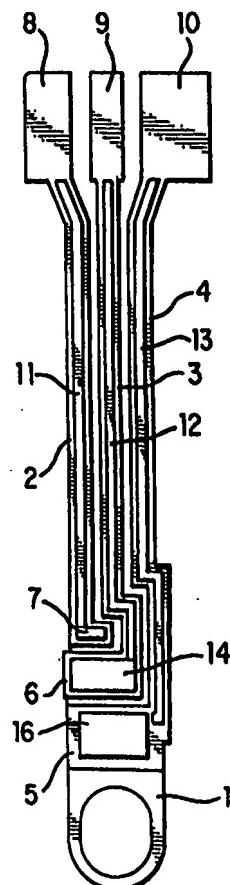
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(71) Applicant (<i>for all designated States except US</i>): ABBOTT LABORATORIES [US/US]; CHAD 0377/AP6D-2, 100 Abbott Park Road, Abbott Park, IL 60064-3500 (US).		
(72) Inventors; and		Published
(75) Inventors/Applicants (<i>for US only</i>): FORROW, Nigel, J. [GB/GB]; 6 Grundy Close, Abingdon, Oxon OX14 3SD (GB). SANGHERA, Gurdial, S. [GB/US]; 4A Crosby Drive, Bedford, MA 01730 (US). WATKIN, Jared, L. [GB/GB]; 11 Kysbie Place, Abingdon, Oxfordshire OX14 1XZ (GB). WALTERS, Stephen [GB/GB]; Flat 4, Stanmore Road, Edgebaston, Birmingham B16 9TB (GB).		<i>With international search report.</i> <i>Before the expiration of the time limit for amending the claims and to be republished in the event of the receipt of amendments.</i>
(74) Agents: POPE, Lawrence, S. et al.; Abbott Laboratories, CHAD 0377/AP6D-2, 100 Abbott Park Road, Abbott Park, IL 60064-3500 (US).		

(54) Title: BIOSENSOR ELECTRODE MEDIATORS FOR REGENERATION OF COFACTORS

(57) Abstract

The present invention is based on the discovery of NAD⁺ and NADP⁺ mediator compounds that do not bind irreversibly to thiol groups in the active sites of intracellular dehydrogenase enzymes. Such mediator compounds avoid a common mode of enzyme inhibition. The mediators can therefore increase the stability and reliability of the electrical response in amperometric electrodes constructed from NAD- or NADP-dependent enzymes.



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BIOSENSOR ELECTRODE MEDIATORS FOR REGENERATION OF COFACTORS

Background of the Invention

The invention is in the general field of electrodes for amperometric biosensors. More specifically, the invention is in the field of compounds for use as mediators for the recycling of cofactors used in these electrodes.

NAD- and NADP-dependent enzymes are of great interest insofar as many have substrates of clinical value, such as glucose, D-3-hydroxybutyrate, lactate, ethanol, and cholesterol. Amperometric electrodes for detection of these substrates and other analytes can be designed by incorporating this class of enzymes and establishing electrical communication with the electrode via the mediated oxidation of the reduced cofactors NADH and NADPH.

NAD- and NADP-dependent enzymes are generally intracellular oxidoreductases (EC 1.x.x.x). The oxidoreductases are further classified according to the identity of the donor group of a substrate upon which they act. For example, oxidoreductases acting on a CH-OH group within a substrate are classified as EC 1.1.x.x whereas those acting on an aldehyde or keto-group of a substrate are classified as EC 1.2.x.x. Some important analytes (e.g., glucose, D-3-hydroxybutyrate, lactate, ethanol, and cholesterol) are substrates of the EC 1.1.x.x enzymes.

The category of oxidoreductases is also broken down according to the type of acceptor utilized by the enzyme. The enzymes of relevance to the present invention have NAD⁺ or NADP⁺ as acceptors, and are classified as EC 1.x.1.x. These enzymes generally possess sulphhydryl groups within their active sites and hence can be irreversibly inhibited by thiol-reactive reagents such as iodoacetate. An irreversible inhibitor forms a stable compound, often through the formation of a covalent bond with a particular amino acid residue (e.g., cysteine, or Cys) that is essential for enzymatic activity. For example, glyceraldehyde-3-P dehydrogenase (EC 1.2.1.9) is stoichiometrically alkylated by iodoacetate at Cys₁₄₉ with concomitant loss of catalytic activity. In addition, the enzymes glucose dehydrogenase, D-3-hydroxybutyrate

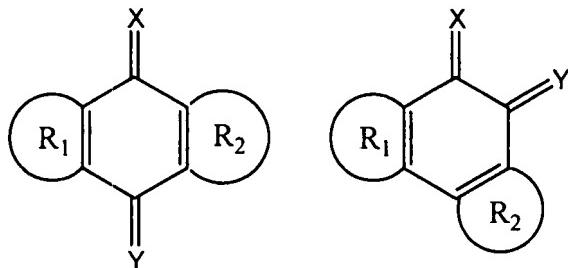
dehydrogenase (HBDH), and lactate dehydrogenase are known to be irreversibly inhibited by thiol reagents. Thus, in seeking to develop stable biosensors containing NAD- or NADP-dependent dehydrogenases, avoidance of compounds that are reactive toward thiols is imperative, as they can act as enzyme inhibitors.

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Summary of the Invention

The present invention is based on the discovery of NAD⁺ and NADP⁺ mediator compounds that do not bind irreversibly to thiol groups in the active sites of intracellular dehydrogenase enzymes. Such mediator compounds avoid a common mode of enzyme inhibition. The mediators can therefore increase the stability and 10 reliability of the electrical response in amperometric electrodes constructed from NAD- or NADP-dependent enzymes.

In one embodiment, the invention features a test element for an amperometric biosensor. The element includes an electrode, which has test reagents distributed on it. The test reagents include a nicotinamide cofactor-dependent 15 enzyme, a nicotinamide cofactor, and a mediator compound having one of the formulae:



or a metal complex or chelate thereof,
where X and Y can independently be oxygen, sulphur, CR³R⁴, NR³, or NR³R⁴⁺; R₁ and
R₂ can independently be a substituted or unsubstituted aromatic or heteroaromatic
group; and R³ and R⁴ can independently be a hydrogen atom, a hydroxyl group or a
5 substituted or unsubstituted alkyl, aryl, heteroaryl, amino, alkoxy, or aryloxy group.
In some cases, either X or Y can be the functional group CZ¹Z², where Z¹ and Z² are
electron withdrawing groups.

- Any alkyl group, unless otherwise specified, may be linear or branched and
may contain up to 12, preferably up to 6, and especially up to 4 carbon atoms.
- 10 Preferred alkyl groups are methyl, ethyl, propyl and butyl. When an alkyl moiety
forms part of another group, for example the alkyl moiety of an alkoxy group, it is
preferred that it contains up to 6, especially to 4, carbon atoms. Preferred alkyl
moieties are methyl and ethyl.

An aromatic or aryl group may be any aromatic hydrocarbon group and may
15 contain from 6 to 24, preferably 6 to 18, more preferably 6 to 16, and especially 6 to
14, carbon atoms. Preferred aryl groups include phenyl, naphthyl, anthryl,
phenanthryl and pyrlyl groups especially a phenyl or naphthyl, and particularly a
phenyl group. When an aryl moiety forms part of another group, for example, the aryl
moiety of an aryloxy group, it is preferred that it is a phenyl, naphthyl, anthryl,
20 phenanthryl or pyrlyl, especially phenyl or naphthyl, and particularly a phenyl, moiety.

A heteroaromatic or heteraryl group may be any aromatic monocyclic or
polycyclic ring system, which contains at least one heteroatom. Preferably, a
heteroaryl group is a 5 to 18-membered, particularly a 5- to 14-membered, and
especially a 5- to 10-membered, aromatic ring system containing at least one
25 heteroatom selected from oxygen, sulphur and nitrogen atoms. 5- and 6-membered
heteroaryl groups, especially 6-membered groups, are particularly preferred.

Heteroaryl groups containing at least one nitrogen atom are especially preferred.
Preferred heteroaryl groups include pyridyl, pyrylium, thiopyrylium, pyrrolyl, furyl,
thienyl, indolinyl, isoindolinyl, indolizinyl, imidazolyl, pyridonyl, pyronyl,

pyrimidinyl, pyrazinyl, oxazolyl, thiazolyl, purinyl, quinolinyl, isoquinolinyl, quinoxalinyl, pyridazinyl, benzofuranyl, benzoxazolyl and acridinyl groups.

When any of the foregoing substituents are designated as being substituted, the substituent groups which may be present may be any one or more of those customarily employed in the development of compounds for use in electrochemical reactions and/or the modification of such compounds to influence their structure/activity, solubility, stability, mediating ability, formal potential (E°) or other property. Specific examples of such substituents include, for example, halogen atoms, oxo, nitro, cyano, hydroxyl, cycloalkyl, alkyl, haloalkyl, alkoxy, haloalkoxy, amino, alkylamino, dialkylamino, formyl, alkoxycarbonyl, carboxyl, alkanoyl, alkylthio, alkylsulphinyl, alkylsulphonyl, arylsulphinyl, arylsulphonyl, carbamoyl, alkylamido, aryl or aryloxy groups. When any of the foregoing substituents represents or contains an alkyl substituent group, this may be linear or branched and may contain up to 12, preferably up to 6, and especially up to 4, carbon atoms. A cycloalkyl group may contain from 3 to 8, preferably from 3 to 6, carbon atoms. An aryl group or moiety may contain from 6 to 10 carbon atoms, phenyl groups being especially preferred. A halogen atom may be a fluorine, chlorine, bromine or iodine atom and any group which contains a halo moiety, such as a haloalkyl group, may thus contain any one or more of these halogen atoms.

An electron withdrawing group may be any group, which forms a stable methylene group CZ^1Z^2 . Such electron withdrawing groups may include halogen atoms, nitro, cyano, formyl, alkanoyl, carboxyl and sulphonic acid groups.

Preferably, X and Y are both oxygen atoms.

It is also preferred that R_1 and R_2 are independently selected from phenyl, naphtuyl, pyridyl and pyrrolyl groups with pyridyl groups being especially preferred. The term "pyridyl group" also includes the N-oxide thereof as well as pyridinium and N-substituted pyridinium groups.

Preferably, R₁ and R₂ are unsubstituted or substituted only by one or more, preferably one or two, alkyl groups, especially methyl groups. It is especially preferred that R₁ and R₂ are unsubstituted.

R₃ and R₄, if present, are preferably independently selected from hydrogen atoms and alkyl groups.

Metal complex and chelates include complexes and chelates with transition metals, especially first-, second- and third-row transition elements such as ruthenium, chromium, cobalt, iron, nickel and rhenium, with ruthenium being particularly preferred. Other groups such as 4-vinyl-4'-methyl-2, 2'-bipridyl (v-bpy) and 10 bipyridyl (bpy) groups may also be included in such complexes and chelates as parts of a complex metal ion. Typically, such complexes and chelates will form as a result of heteroatoms in R₁ and R₂ coordinating with a metal ion or metal ion complex.

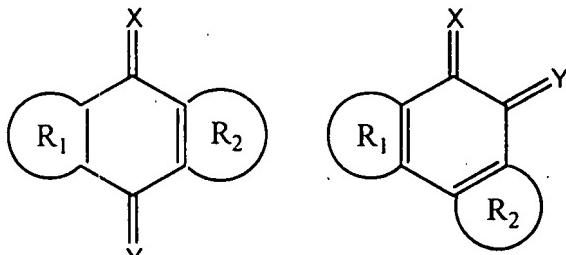
The test reagents can be deposited on the electrode in one or more ink-based layers. The test reagents can be screen-printed onto the working electrode in a single 15 layer.

The element can be an amperometric dry-strip sensor that includes an elongated, electrically insulating carrier having a pair of longitudinal, substantially parallel electrically conducting tracks thereupon, and a pair of electrodes. The electrodes can each be electrically connected to a different one of the tracks; one of 20 the electrodes can be a reference/counter electrode, while another electrode can be a working electrode. The element can also include a dummy electrode. Further, the element can include a membrane positioned to filter samples prior to their introduction onto the electrodes.

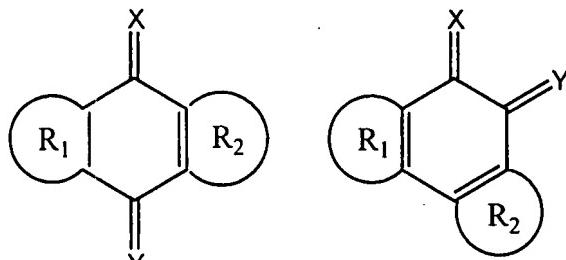
The sensor can additionally include a supporting strip of electrically insulating 25 carrier material (e.g., a synthetic polymer such as polyvinyl chloride, or a blend of synthetic polymers).

The mediator compound can be a quinone. Examples of suitable quinones include 1,10-phenanthroline quinone, 1,7-phenanthroline quinone, and 4,7-phenanthroline quinone.

In another embodiment, the invention features an electrode strip for an amperometric sensor having a readout. The strip includes a support adapted for releasable attachment to the readout, a first conductor extending along the support and comprising a conductive element for connection to the readout; a working electrode in contact with the first conductor and positioned to contact a sample mixture; a second conductor extending along the support and comprising a conductive element for connection to the readout; and a reference/counter electrode in contact with the second conductor and positioned to contact the sample and the second conductor. The active electrode of the strip includes a mediator compound having one of the formulae:

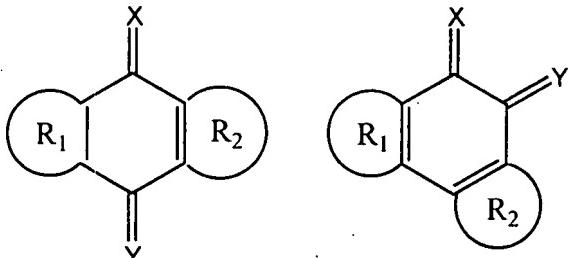


Still another embodiment of the invention features a method for mediating electron transfer between an electrode and a nicotinamide cofactor. The method includes the steps of using a mediator compound in the presence of a nicotinamide cofactor-dependent enzyme, where the mediator compound is a quinoid compound that is incapable of binding irreversibly to the thiol groups. The mediator compound can, for example, have reactive unsaturated bonds in adjacent aromatic ring. Suitable mediator compounds include those having the formulae:



For example, the mediator compound can be 1,10-phenanthroline quinone, 1,7-phenanthroline quinone, or 4,7-phenanthroline quinone.

In yet another embodiment, the invention features a printing ink. The ink includes a nicotinamide cofactor-dependent enzyme, a nicotinamide cofactor, and a mediator compound having one of the formulae:



- For example, the mediator compound can be 1,10-phenanthroline quinone,
5 1,7-phenanthroline quinone, or 4,7-phenanthroline quinone. The enzyme can be, for example, alcohol dehydrogenase, lactate dehydrogenase, 3-hydroxybutyrate dehydrogenase, glucose-6-phosphate dehydrogenase, glucose dehydrogenase, formaldehyde dehydrogenase, malate dehydrogenase, or 3-hydroxysteroid dehydrogenase.

10 Unless otherwise defined, all technical and scientific terms used herein have the same meaning as commonly understood by one of ordinary skill in the art to which this invention belongs. Although methods and materials similar or equivalent to those described herein can be used in the practice or testing of the present invention, the preferred methods and materials are described below. All publications, 15 patent applications, patents, technical manuals, and other references mentioned herein are incorporated by reference in their entirety. In case of conflict, the present application, including definitions, will control. In addition, the materials, methods, and examples are illustrative only and not intended to be limiting.

An advantage of the new mediators is their non-reactivity with respect to 20 active-site thiol groups in enzymes. This improves the stability and the shelf life of biosensor electrodes to an unexpected degree. Also as a result of this stability, the enzyme and mediator can be incorporated together in a printing ink or dosing solution to facilitate construction of the biosensors. The use of a mediator that is not an irreversible inhibitor of the enzyme will result in the retention of a large proportion of

enzyme activity during the biosensor manufacture. NAD- and NADP-dependent dehydrogenase enzymes are generally expensive and labile and improvement of their stability is therefore highly desirable.

Advantageously, the compounds disclosed herein can also be used as
5 mediators to the cofactors NADH and NADPH coupled with a wide range of NAD- or NADP-dependent enzymes; as labels for antigens or antibodies in immunochemical procedures; and in other applications in the field of electrochemistry and
bioelectrochemistry. The mediators require low oxidation potentials for re-oxidation
following the reaction with NADH or NADPH. This is of particular advantage when
10 testing in whole blood, in which the potential for interference from exogenous
electroactive species (e.g., ascorbic acid, uric acid) is particularly high. The low
potential can be advantageous because it can obviate the need for a dummy electrode
to remove electroactive species in the sample. Also, the oxidized native form of the
mediator can decrease the background current that would be present with a reduced
15 mediator.

Other features and advantages of the invention will be apparent from the
following detailed description.

Brief Description of the Drawings

Fig. 1 is an exploded view of an electrode strip according to one embodiment
20 of the invention.

Fig. 2 is a representation of an assembled electrode strip.

Fig. 3 is a graphical plot of current in μA against NADH concentration in mM
for printed electrodes containing 1,10-phenanthroline quinone.

Fig. 4 is a graphical plot of current in μA against NADH concentration in mM
25 for printed electrodes containing Meldola's Blue.

Fig. 5 is a bar chart displaying residual enzyme activity (i.e., as a percentage of
the initial activity) after incubation of HBDH with various mediators.

Fig. 6 is a graphical plot of current in μ A against D-3-hydroxybutyrate concentration in mM for printed electrodes containing 1,10-phenanthroline quinone, D-3-hydroxybutyrate dehydrogenase and NAD⁺ tested after 4, 14 and 26 weeks.

Fig. 7 is a graphical plot of current in μ A against D-3-hydroxybutyrate concentration in mM for printed electrodes containing Meldola's Blue, D-3-hydroxybutyrate dehydrogenase, and NAD⁺ tested after 2 and 14 weeks, respectively.

Fig. 8 is a graphical plot of calibrated response to glucose in whole blood for printed electrodes containing 1,10-phenanthroline quinone, glucose dehydrogenase, and NAD⁺.

Detailed Description of the Invention

A class of compounds, selected for their inability to combine irreversibly with thiols, are disclosed for use as NADH or NADPH mediators. The structural, electronic, and steric characteristics of these mediators render them nearly incapable of reacting with thiols. Because these mediators are virtually precluded from binding irreversibly to the active site sulphhydryl groups of NAD- and NADP-dependent dehydrogenases, inactivation of the enzyme and consequent loss of biosensor stability is circumvented.

The NADH and NADPH mediators can be used in the manufacture of amperometric enzyme sensors for an analyte, where the analyte is a substrate of an NAD- or NADP-dependent enzyme present in the sensor, such as those of the kind described in EP 125867-A. Accordingly, amperometric enzyme sensors of use in assaying for the presence of an analyte in a sample, especially an aqueous sample, can be made. For example, the sample can be a complex biological sample such as a biological fluid (e.g., whole blood, plasma, or serum) and the analyte can be a naturally occurring metabolite (e.g., glucose, D-3-hydroxybutyrate, ethanol, lactate, or cholesterol) or an introduced substance such as a drug.

Of particular utility for the manufacture of amperometric enzyme sensors, the present invention further provides an ink that includes the NADH and NADPH mediators disclosed herein.

The present invention also includes any precursor, adduct, or reduced (leuco) form of the above mediators that can be converted *in situ* by oxidation or decomposition to the corresponding active mediators. Such precursors or adducts can include hemiacetals, hemithioacetals, cyclic acetals, metal o-quinone complexes, protonated forms, acetone adducts, etc.

A non-limiting list of enzymes that can be used in conjunction with the new mediators is provided in Table 1.

TABLE 1

1.1.1.1	Alcohol Dehydrogenase
1.1.1.27	Lactate Dehydrogenase
1.1.1.31	3-Hydroxybutyrate Dehydrogenase
1.1.1.49	Glucose-6-phosphate Dehydrogenase
1.1.1.47	Glucose Dehydrogenase
1.2.1.46	Formaldehyde Dehydrogenase
1.1.1.37	Malate Dehydrogenase
1.1.1.209	3 -hydroxysteroid Dehydrogenase

Amperometric enzyme sensors adopting the mediators of the present invention generally use a test element, for example, a single-use strip. A disposable test element can carry a working electrode, for example, with the test reagents including the enzyme, the nicotinamide cofactor (i.e., NAD⁺ or NADP⁺), the mediators of the present invention for generation of a current indicative of the level of analyte, and a

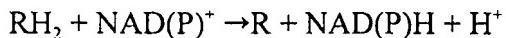
reference/counter electrode. The test reagents can be in one or more ink-based layers associated with the working electrode in the test element. Accordingly, the sensor electrodes can, for example, include an electrode area formed by printing, spraying, or other suitable deposition technique.

5 Referring to Figs. 1 and 2, an electrode support 1, typically made of PVC, polycarbonate, or polyester, or a mixture of polymers (e.g., Valox, a mixture of polycarbonate and polyester) supports three printed tracks of electrically conducting carbon ink 2, 3, and 4. The printed tracks define the position of the working electrode 5 onto which the working electrode ink 16 is deposited, the reference/counter 10 electrode 6, the fill indicator electrode 7, and contacts 8, 9, and 10.

The elongated portions of the conductive tracks are respectively overlaid with silver/silver chloride particle tracks 11, 12, and 13 (with the enlarged exposed area 14 of track 12 overlying the reference electrode 6), and further overlaid with a layer of hydrophobic electrically insulating material 15 that leaves exposed only positions of 15 the reference/counter electrode 14, the working electrode 5, the fill indicator electrode 7, and the contact areas 8, 9, and 10. This hydrophobic insulating material serves to prevent short circuits. Because this insulating material is hydrophobic, it can serve to confine the sample to the exposed electrodes. A suitable insulating material is Sericard, commercially available from Sericol, Ltd. (Broadstairs, Kent, UK). 20 Optionally, a first mesh layer 17, a second insulative layer 18, a second mesh layer 19, a third insulative layer 20, and a tape 21 can overlay the hydrophobic insulating material.

Respective ink mixtures can be applied onto a conductive track on a carrier, for example, in close proximity to a reference electrode 14 connected to a second 25 track. In this way, a sensor can be produced, which is capable of functioning with a small sample of blood or other liquid covering the effective electrode area 5. The mixtures are preferably, but not exclusively, applied to the carrier by screen printing.

In general, NAD(P)-dependent dehydrogenases catalyze reactions according to the equation:



where RH_2 represents the substrate (analyte) and R the product. In the process of the forward reaction, NAD(P)^+ (i.e., NAD^+ or NADP^+) is reduced to NAD(P)H . Suitable amperometric biosensors provide an electrochemical mediator that can reoxidize

- 5 NAD(P)H, thereby regenerating NAD(P)^+ . Reoxidation occurs at an electrode to generate a current that is indicative of the concentration of the substrate.

In one embodiment, a dry sensor is provided. The sensor includes an elongated electrically insulating carrier having a pair of longitudinal, substantially parallel, electrically conducting tracks thereupon, each track being provided at the same end with means for electrical connection to a read-out and provided with an electrode, one of the electrodes being the reference/counter electrode and the other being the working electrode, together with test reagents. The sensor can be configured in the form of a supporting strip of electrically insulating carrier material such as a synthetic polymer (e.g., PVC, polycarbonate, or polyester, or a mixture of polymers such as Valox) carrying the two electrodes supported on electrically conductive tracks between its ends. For example, the electrodes can take the form of two rectangular areas side by side on the carrier strip, as shown in Fig. 2 (i.e., electrodes 14 and 16). Such areas can be designed as a target area to be covered by a single drop of sample, such as whole blood, for testing the analyte. If desired, non-rectangular areas (e.g., diamond-shaped, semicircular, circular, or triangular areas) can be employed to provide a target area for optimized contact by a liquid sample.

The carrier includes at least two electrodes, namely a reference/counter electrode and a working electrode. Other electrodes such as a dummy electrode can also be included. These other electrodes can be of similar formulation to the working electrode (i.e., with the associated test reagents), but lacking one or more of the working electrode's active components. A dummy electrode, for example, can provide more reliable results, in that if charge passed at the dummy electrode is

subtracted from charge passed at the working electrode, then the resulting charge can be concluded to be due to the reaction of interest.

A membrane can be provided at or above the target to perform a filtration function. For example, a membrane can filter blood cells from a sample before the 5 sample enters the test strip. Examples of commercially available membranes that can be used include Hemasep V, Cytosep, and Hemadyne (Pall Biosupport, Fort Washington, NY 11050). As an alternative, a filtration or cellular separation membrane can be cast *in situ*. This can be achieved by casting hydrophobic polymers such as cellulose acetate, polyvinyl butyral and polystyrene and/or hydrophilic 10 polymers such as hydroxypropyl cellulose, polyvinylpyrrolidone, polyvinyl alcohol and polyvinyl acetate.

In another embodiment, there is provided a single use disposable electrode strip for attachment to signal readout circuitry of a sensor system. The strip can detect a current representative of an analyte in a liquid mixture. The strip includes an 15 elongated support adapted for releasable attachment to the readout circuitry; a first conductor extending along the support and including a conductive element for connection to the readout circuitry; a working electrode on the strip in contact with the first conductor and positioned to contact the mixture; a second conductor extending along the support, comprising a conductive element for connection to the readout 20 circuitry; and a reference/counter electrode in contact with the second conductive element and positioned to contact the mixture and the second conductor as depicted in Fig. 1.

The working electrode can include a printed layer on the support, and the 25 printed layer itself can include an NAD or NADP-dependent dehydrogenase enzyme capable of catalyzing a reaction involving a substrate for the enzyme. This layer can also include the corresponding nicotinamide cofactor and a mediator of the present invention capable of transferring electrons between the enzyme-catalyzed reaction and the first conductor via NADH or NADPH, to create a current representative of the activity of both the enzyme and the analyte.

The first conductive element and the active electrode can be spaced apart from the second conductive element and the reference/counter electrode, and the electrodes sized and positioned to present a combined effective area small enough to be completely covered by a drop of blood or other test sample; typically the reaction 5 zone is 5 mm² but can be as large as 25 mm². The test sample completes an electrical circuit across the active electrode and the reference/counter electrode for amperometric detection of the activity of the enzyme.

In a preferred embodiment of the present invention a working electrode is produced by using a formulation which includes not only the enzyme, nicotinamide 10 cofactor and the mediator but also filler and binder ingredients which cause the working electrode to give an increasing monotonic response to concentrations of interest for the analyte being sensed when measured in a kinetic mode in which oxidation and reduction of the mediator both occur during the measurement. The concept is to provide a stable reaction layer on the surface of the working electrode 15 when the sample is applied. This allows the use of mediators which are sparingly soluble in the sample. As the mediator is reduced by reaction with the enzyme, cofactor and analyte, it is retained in close proximity to the electrode surface so that it can be readily reoxidized without significant loss to precipitation. The maintenance of this thin reaction layer also allows the overall analytical reaction to occur in a small 20 volume of the overall sample so in effect what is measured is the flux of analyte from the bulk specimen to this reaction layer.

This reaction layer needs to remain stable for at least the time to conduct a reproducible kinetic measurement. Typical times for such a measurement range between about 5 and 60 seconds, although stability for longer times are preferred. 25 Typically, the disposable electrode strips of interest are mass produced and therefore it is desirable to have a safety margin with regard to any required property to account for the inherent variability in any mass manufacturing process.

The stability of the reaction layer can be improved by a proper combination of fillers and binders. The layer is preferably sufficiently stable to give an approximately

linear reproducible response in a kinetic measurement over the concentration range of interest for a given analyte. For instance, for Ketone bodies (measured as hydroxybutyrate) this would be between about 1 and 8 mM while for glucose it would be between about 2 and 40 mM.

5 The kinetic measurement involves the cycling of the mediator between an oxidized state and a reduced state. The rate of this cycling, which is reflected in the current observed during the course of the test, is dependent upon the concentration of the analyte in the sample. The greater the concentration of the analyte the more enzyme cofactor which is reduced in the course of the enzyme oxidizing the analyte.

10 The mediator in turn becomes reduced in reoxidizing the cofactor and is then reoxidized at the electrode surface. However, because of its very low solubility only a small amount of mediator is immediately available to react with the reduced cofactor. Consequently mediator which reacts with reduced cofactor and is reoxidized at the electrode will then react with further reduced cofactor and this continues through the

15 course of a kinetic measurement. Thus the greater the concentration of the reduced cofactor (reflective of a greater concentration of analyte in the sample) the greater the driving force for the cycling of the mediator and thus the greater the rate of cycling.

In some cases the cofactor may also engage in cycling between an oxidized state and a reduced state during the kinetic measurement. This depends upon whether 20 there is a sufficient quantity of cofactor initially present to convert all the analyte present in the reaction layer. If there is insufficient cofactor initially present as oxidized cofactor is regenerated it promotes the oxidation of any analyte remaining in the reaction layer by becoming reduced again.

However, what is critical is that a given concentration of analyte reproducible 25 results in the production of the same signal in the kinetic test for a particular electrode strip design and that the signal increases monotonically, preferably linearly, with the concentration of the analyte (in other words that the signal be a true function of the analyte concentration) over the concentration range of interest. This allows the manufacturer of the electrode strips to establish a universal calibration for a given lot

of electrode strips such that any given signal obtained from a given strip under standard test conditions uniquely correlates to a particular analyte concentration. Thus it is important that within the concentration range of interest there be no uncontrollable variable other than the analyte concentration which would substantially effect the signal.

The signal may be the current observed at a fixed time after the test is initiated or it may be the current integrated over some period occurring some fixed time after the test is initiated (in essence the charge transferred over some such period). The test is conducted by covering the working electrode and a reference/counter electrode with sample and then applying a potential between them. The current which then flows is observed over some time period. The potential may be imposed as soon as the sample covers the electrodes or it may be imposed after a short delay, typically about 3 seconds, to ensure good wetting of the electrodes by the sample. The fixed time until the current or current integration is taken as the signal should be long enough to ensure that the major variable affecting the observed current is the analyte concentration.

The reference electrode/counter electrode may be a classic silver/silver chloride electrode but it may also be identical to the working electrode in construction. In one embodiment the two separate conductive tracks may both be coated with an appropriate formulation of enzyme, cofactor and mediator in a binder and filler containing aqueous vehicle to yield a coating. In those cases in which the coating is non-conductive, e.g. when the filler is a non-conductor, a common coating may overlay both electrodes. When a potential is applied one of the electrodes will function as a reference/counter electrode by absorbing the electrons liberated at the other, working, electrode. The mediator at the reference/counter electrode will simply become reduced as a result of interaction with the electron flow at its electrode.

The reaction layer which yields the desired behavior is obtained by formulating the working electrode with binder and filler ingredients. The object is to allow the sample to interact with the enzyme, cofactor and mediator but to also ensure

that these chemically active ingredients remain in the immediate vicinity of the surface of the electrode. The binder ingredient should include materials which readily increase the viscosity of aqueous media and promote the formation of films or layers. Typical of such materials are the polysaccharides such as guar gum, alginate, locust 5 bean gum, carrageenan and xanthan. Also helpful are materials commonly known as film formers such as polyvinyl alcohol (PVA), polyvinyl pyrrole, cellulose acetate, carboxymethyl cellulose and poly (vinyl oxazolidinone). The filler ingredient should be a particulate material which is chemically inert to the oxidation reduction reactions involved in the measurement and insoluble in aqueous media. It may be electrically 10 conductive or non-conductive. Typical materials include carbon, commonly in the form of graphite, titanium dioxide, silica and alumina.

The active electrode may be conveniently produced by formulating the enzyme, cofactor, mediator and binder and filler ingredients into an aqueous vehicle and applying it to the elongated, electrically insulating carrier having conducting 15 tracks. The formulation may be applied by printing such as screen printing or other suitable techniques. The formulation may also include other ingredients such as a buffer to protect the enzyme during processing, a protein stabilizer to protect the enzyme against denaturation and a defoaming agent. These additional ingredients may also have an effect on the properties of the reaction layer.

20 The working electrode typically has a dry thickness between about 2 and 50 microns preferably between about 10 and 25 microns. The actual dry thickness will to some extent depend upon the application technique used to apply the ingredients which make up the working electrode. For instance thicknesses between about 10 and 25 microns are typical for screen printing.

25 However, the thickness of the reaction layer is not solely a function of the dry thickness of the working electrode but also depends upon the effect of the sample on the working electrode. In the case of aqueous samples the formulation of the working electrode ingredients will effect the degree of water uptake this layer displays.

The filler typically makes up between about 20 and 30 weight percent of the aqueous vehicle. The amounts of the other ingredients are typically less than about 1 weight percent of the aqueous vehicle and are adjusted empirically to achieve the desired end properties. For instance, the amount of buffer and protein stabilizer are 5 adjusted to achieve the desired degree of residual enzyme activity. In this regard one may use more enzyme and less stabilizer or less enzyme and more stabilizer to achieve the same final level of enzyme activity. The amount of binder and defoaming agent should be adjusted to give suitable viscosities for the method of application with higher viscosities being suitable for screen printing and lower viscosities being 10 suitable for rotogravure printing.

A suitable aqueous ink formulation can be formulated in accordance with Table 2 with the balance being deformer, buffer, enzyme activity enhancers and water to make up 1 gram of formulated ink.

TABLE 2

Enzyme (such as Glucose Dehydrogenase or 3-hydroxybutyrate Dehydrogenase)	200 to 4000 Units
Nicotinamide cofactor (such as NAD)	5 to 30 weight percent
Mediator (such as 1, 10 phenanthroline quinone)	0.1 to 1.5 weight percent
Filler (such as ultra fine carbon or titania)	10 to 30 weight percent
Binder (such as alginate or guar gum)	0.01 to 0.5 weight percent
Protein stabilizer (such as Trehalose or Bovine Serum Albumin)	0.01 to 2 weight percent

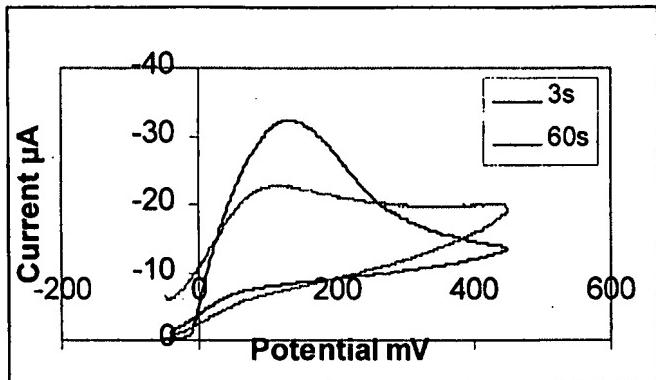
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The stability of the reaction layer can be readily evaluated using cyclic voltammetry with various time delays. The working electrode formulation is evaluated by exposing it to a sample containing a relatively high concentration of analyte and subjecting it to a steadily increasing potential to a maximum value and 20 then a steadily decreasing potential back to no applied potential. The resulting current

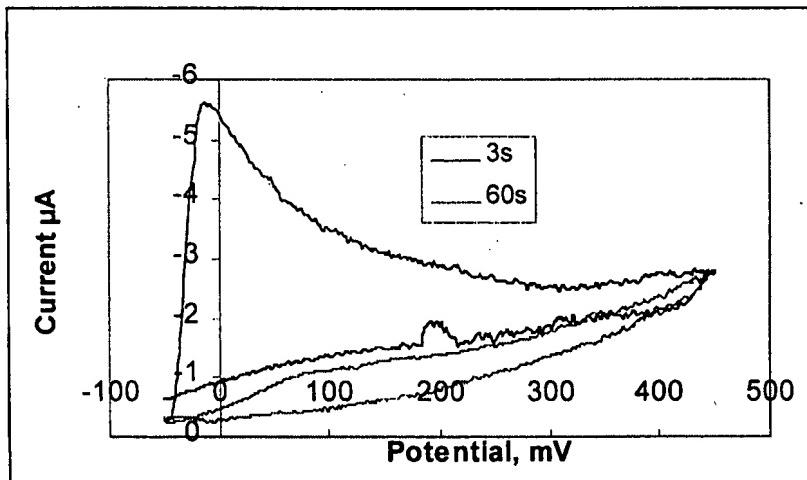
increases to a peak value and then drops off as the voltage sweep continues. Such cyclic voltammetry evaluations are conducted after various delay periods after the working electrode is exposed to the sample. The change in peak current with increasingly long delay periods is a measure of the stability of the reaction layer. The 5 more stable the reaction layer the smaller the decrease in peak current.

An evaluation was conducted to compare the stability of a working electrode formulated in accordance with the teachings of the present invention to that of a "working electrode" formulated according to the teachings of Geng et al. at pages 1267 to 1275 of Biosensors and Bioelectronics, Volume II, number 12 (1996). The 10 working electrode representative of the present invention was formulated with about 25 weight percent filler (ultra fine carbon), binder, protein stabilizer and deformers as taught hereinabove and the working electrode representative of Geng was formulated with a high molecular weight poly (ethylene oxide) as described at page 1267 of the Geng article. In each case a potential was applied at a scan rate of 50 millivolt per 15 second up to 400 mV versus a silver/silver chloride reference electrode after exposing the working electrode to a 20 mM aqueous solution of glucose for 3 seconds and 60 seconds. The formulation according to the present invention yields a stable reaction layer in which the peak current after 60 seconds is 60% of that observed after 3 seconds while the formulation according to the Geng article yields an unstable 20 reaction layer in which no peak current is observable after 60 seconds exposure.

This is attributed to a dissolution of the electrode with a loss of the reagents to the bulk solution. The respective voltammograms are as follows:



Formulation with Binder and Filler (Present Invention)



5

PEO formulation (Geng Paper)

The test strips of this invention can detect analytes that are substrates of NAD-
or NADP-dependent dehydrogenase enzymes using a mediator selected from the
10 compounds disclosed herein, such as 1,10-PQ.

Test strips according to this invention are intended for use with electronic
apparatus and meter systems. These control the progress of the electrochemical
reaction (e.g., by maintaining a particular potential at the electrodes), monitor the
reaction, and calculate and present the result. A particular feature that is desirable in a
15 meter system for use with test strips of this type is the capability of detecting the
wetting of the reaction zone by sample fluid, thus allowing timely initiation of the

measurement and reducing the potential for inaccuracies caused by user error. This goal can be achieved by applying a potential to the electrodes of the test strip as soon as the strip is inserted into the meter; this potential can be removed for a short time to allow wetting to be completed before initiation of measurement.

5 The meter can also feature a means for automatically identifying test strips for measuring different analytes. This can be achieved, for example, when one or more circuit loops are printed on the test strip; each loop can provide a resistance characteristic of the type of strip, as described in U.S. Pat. No. 5,126,034 at column 4, lines 3 to 17. As a further alternative, notches or other shapes might be cut into the
10 proximal end of the test strip; switches or optical detectors in the meter can detect the presence or absence of each notch. Other strip-type recognition techniques include varying the color of the strips and providing the meter with a photodetector capable of distinguishing the range of colors; and providing the strips with barcodes, magnetic strips, or other markings, and providing the meter with a suitable reading
15 arrangement.

In one example of a test strip for large scale production, the strip electrodes have a two-electrode configuration comprising a reference/counter electrode and a working electrode. The carrier can be made from any material that has an electrically insulating surface, including poly(vinyl chloride), polycarbonate, polyester, paper,
20 cardboard, ceramic, ceramic-coated metal, blends of these materials (e.g., a blend of polycarbonate and polyester), or another insulating substance.

A conductive ink is applied to the carrier by a deposition method such as screen printing. This layer forms the contact areas, which allow the meter to interface with the test strip, and provides an electrical circuit between the contacts and the
25 active chemistry occurring on the strip. The ink can be an air-dried, organic-based carbon mixture, for example. Alternative formulations include water-based carbon inks and metal inks such as silver, gold, platinum, and palladium. Other methods of drying or curing the inks include the use of infrared, ultraviolet, and radio-frequency radiation.

A layer forming the reference/counter electrode is printed with an organic solvent-based ink containing a silver/silver chloride mixture. Alternative reference couples include Ag/AgBr, Ag/AgI, and Ag/Ag₂O. The print extends to partially cover the middle track of the carbon print where it extends into the reaction zone. It is 5 useful if separate parts of this print are extended to cover parts of other carbon tracks outside the reaction zone, so that the total electrical resistance of each track is reduced.

A layer of dielectric ink can optionally be printed to cover the majority of the printed carbon and silver/silver chloride layers. In this case, two areas are left 10 uncovered, namely the electrical contact areas and the sensing area which will underlie the reactive zone as depicted in Figs. 1 and 2. This print serves to define the area of the reactive zone, and to protect exposed tracks from short circuit.

For the working electrode, one or more inks are deposited to a precise thickness within a defined area on top of one of the conductive tracks within the 15 reaction zone, to deposit the enzyme, cofactor and a mediator of the present invention. It is convenient to do this by means of screen printing. Other ways of laying down this ink include inkjet printing, volumetric dosing, gravure printing, flexographic printing, and letterpress printing. Optionally, a second partially active ink can be deposited on a second conductive track to form a dummy electrode.

20 Polysaccharides can optionally be included in the ink formulation. Suitable polysaccharides include guar gum, alginate, locust bean gum, carrageenan and xanthan. The ink can also include a film former; suitable film-forming polymers include polyvinyl alcohol (PVA), polyvinyl pyrrole, cellulose acetate, CMC, and poly(vinyl oxazolidinone). Ink fillers can include titanium dioxide, silica, alumina, or 25 carbon.

The following are illustrative, non-limiting examples of the practice of the invention:

EXAMPLE 1

Mediators:

Meldola's Blue (MB) (Compound 3) was obtained as the hemi-ZnCl₂ salt from Polysciences, Inc. 2,6-Dichloroindophenol (DCIP) (Compound 6) and Tris buffer were purchased from Sigma. The phosphate buffered saline (PBS) solution (Dulbecco's formula) was prepared from tablets supplied by ICN Biomedicals, Ltd.

5 D-3-Hydroxybutyrate dehydrogenase (HBDH; EC 1.1.1.30) from *Pseudomonas sp.* was purchased from Toyobo Co., Ltd. p-Nicotinamide adenine dinucleotide (NAD⁺) and D,L-3-hydroxybutyric acid were supplied by Boehringer Mannheim.

10 1,10-Phenanthroline quinone (1,10-PQ) (Compound 7) was prepared according to the method of Gillard *et al.* (*J. Chem. Soc. A*, 1447-1451, 1970). 1,7-Phenanthroline quinone (1,7-PQ) (Compound 8) was synthesized using the procedure described by Eckert *et al.* (*Proc. Natl. Acad. Sci. USA*, 79:2533-2536, 1982). 2,9-Dimethyl-1,10-phenanthroline quinone (2,9-Me₂-1,10-PQ) (Compound 10) was synthesized as a byproduct of the nitration of neocuproine as disclosed by 15 Mullins *et al.* (*J. Chem. Soc., Perkin Trans. I*, 75-81, 1996). 1-Methoxy phenazine methosulphate (1-MeO-PMS) (Compound 5) was prepared via the methylation of 1-methoxy phenazine adapted from the method described by Surrey (*Org. Synth. Coll. Vol. 3*, Ed. E. C. Horning, Wiley, New York, 753-756). 1-Methoxy phenazine was synthesized by a modified Wohl-Aue reaction as reported by Yoshioka (*Yakugaku Zasshi*, 73:23-25, 1953). 4-Methyl-1,2-benzoquinone (4-Me-BQ) (Compound 4) was prepared via oxidation of 4-methyl catechol with o-chloranil according to a general procedure by Carlson *et al.* (*J. Am. Chem. Soc.*, 107:479-485, 1985). The 1,10-PQ complex [Ru(bpy)₂(1,10-PQ)](PF₆)₂ (Compound 12) was obtained from [Ru(bpy)₂Cl₂] (Strem Chemicals, Inc.) as reported by Goss *et al.* (*Inorg. Chem.*, 24:4263-4267, 20 25 1985).

Preparation of 1-Me-1,10-phenanthrolinium quinone trifluoromethane sulphonate (1-Me-1,10-PQ⁺) (Compound 11):

Methyl trifluoromethane sulphonate (Aldrich) (1.0 ml) was added to a solution of 1,10-PQ (0.50 g, 2.38 mmol) in anhydrous methylene chloride (25 ml) under

nitrogen. Immediate precipitation occurred and the resulting mixture was stirred for 24 hours. Filtration followed by washing with methylene chloride afforded 1-Me-1,10-PQ⁺ (0.65 g, 73%) as a fine yellow powder.

Evaluation of Meldola's Blue and 1,10-PQ as NADH Mediators in Dry Strips:

5 Screen-printed electrodes incorporating 1,10-PQ and MB were produced from an organic carbon ink containing these NAD(P)H mediators at a level of 3.5 mg/g ink. The solid mediators were mixed into a commercial conducting carbon ink (Gwent Electronic Materials).

10 The dose response curve for the electrodes containing 1,10-PQ tested with aqueous NADH solutions (0-16.7 mM) in PBS at a poise potential of +400 mV versus a printed Ag/AgCl reference electrode is shown in Fig. 3. A slope of 0.58 μ A mM⁻¹ NADH was recorded. The dose response curve for the electrodes containing MB tested with aqueous NADH solutions (0-12.4 mM) at a poise potential of +100 mV versus a printed Ag/AgCl reference electrode is shown in Fig. 4. An increased slope 15 of 8.48 μ A mM⁻¹ NADH was observed.

Assessment of Mediator Inhibition of D-3-Hydroxybutyrate Dehydrogenase:

20 A series of 18 solutions (2.5 ml each) were prepared, each containing 50 U/ml HBDH and 1.29 or 2.58 mg of the following NAD(P)H mediators: MB(3), 4-Me-BQ(4), 1-MeO-PMS(5), DCIP(6), 1,10-PQ(7), 1,7-PQ(8), 2,9-Me₂-1,10-PQ(10), 1-Me-1,10-PQ⁺(11), and [Ru(bpy)₂(1,10-PQ)](PF₆)₂ (12) in Tris buffer (50 mM, pH 8.2). A control solution was also prepared, containing enzyme but no mediator. The 25 solutions were incubated for 0.5 hours at 37.5 C, then assayed (in triplicate) for NADH at 340 nm, using a Sigma Diagnostics D-3-hydroxybutyrate kit. The extent of the interference of the added mediator with the assay rate compared to the control afforded a quantitative measure of the mediator's efficiency as an oxidant of NADH.

The enzyme was then reisolated from the mediator solutions by filtration through a polysulfone membrane (nominal molecular weight cut-off: 30,000) in a microcentrifuge filter (Millipore). The enzyme remaining on the filter was dissolved in Tris buffer (0.2 ml), and the resulting solution was assayed (in triplicate) with the

Sigma kit. By comparing the results of the assays before and after filtration, the effect of any covalently and/or irreversibly bound mediator on the enzyme activity could be determined.

The results of the two assays on each solution before and after filtration are
5 collected in Table 3.

TABLE 3

Mediator (Compound No.)	Assay Rate (absorbance units/min)		
	control (no mediator)	before filtration	after filtration
1,10-PQ	0.167	0.149	0.160(96%)
1,7-PQ	0.155	0.115	0.150(97%)
MB	0.167	0.008	0.026(16%)
4-Me-BQ	0.170	0.005	0.007(4%)
1-MeO-PMS	0.150	0.009	0.071 (47%)
DCIP	0.150	0.104	0.085 (57%)
2,9-Me ₂ -1,10-PQ	0.197	0.189	n/a
1-Me-1, 10-PQ*	0.197	0.150	0.185 (94%)
[Ru(bpy) ₂ (1,10-PQ)](PF ₆) ₂	0.197	0.114	0.193 (98%)

Although these results demonstrated that the phenanthroline quinone mediators were relatively inefficient NADH mediators compared to Meldola's Blue and 1-MeO-PMS (i.e., the assay rate "before filtration" was depressed only to a small extent), over 90% of the original enzyme activity for the solutions containing 1,10-PQ, 1,7-PQ, 1-MeO-1,10-PQ, or [Ru(bpy)₂(1,10-PQ)](PF₆)₂ was restored "after filtration." This was not the case for MB, 1-MeO-PMS, DCIP, or 4-Me-BQ. Indeed, the quinone mediator 4-Me-BQ proved to be the most potent inhibitor with only 4% of the original activity remaining "after filtration." Thus, the latter four mediators

partially inactivate HBDH while the newly described mediators advantageously had little or no effect on enzyme activity.

The percentage residual enzyme activities for each mediator are displayed as a bar chart in Fig. 5, which reveals that the mediators of the present invention, represented by black bars, are not strong inhibitors of HBDH. In contrast, MB, 4-Me-BQ, 1-MeO-PMS, and DCIP all irreversibly inhibited HBDH, with concomitant losses in activity ranging from 43 to 96%; these results are represented by grey bars in Figure 5.

EXAMPLE 2

10 Evaluation of Meldola's Blue and 1,10-PQ in Dry Strips containing HBDH:

Screen-printed electrodes were produced from an aqueous carbon ink incorporating 1,10-PQ or MB at a level of 2.4 or 4.3 mg/g ink, respectively, together with the enzyme HBDH (120 units/g ink) and NAD⁺ (110 mg/g ink). The ink also contained a polysaccharide binder.

15 The dose response curves for the electrodes containing 1,10-PQ are given in Fig. 6. The electrodes were tested after 4, 14, and 26 weeks of storage (30°C, desiccated) with aqueous D-3-hydroxybutyrate solutions (0-25 mM) in PBS at a poise potential of +400 mV versus a printed Ag/AgCl reference electrode. All three dose responses were non-linear and levelled out with a current of 8.5 μ A being recorded at
20 24 mM D-3-hydroxybutyrate. This demonstrated that the response of the dry electrodes was stable for at least 26 weeks.

The dose response curves for the electrodes containing MB are provided in Fig. 7. The electrodes were tested after 2 and 14 weeks storage (30°C, desiccated) with aqueous D-3-hydroxybutyrate solutions (0-28 mM) in PBS at a poise potential of
25 +100 mV versus a printed Ag/AgCl reference electrode. The dose response curves were similar to those in Fig. 4. A current of 8.6 μ A was recorded at 24 mM D-3-hydroxybutyrate for these electrodes after 2 weeks storage. This is almost identical to responses obtained from dry strips containing 1,10-PQ.

This result demonstrated that the ability of a compound such as MB to mediate very efficiently with NADH compared to 1,10-PQ is outweighed by the fact that it inhibits HBDH. Furthermore, the stability of the electrode response to D-3-hydroxybutyrate is compromised through the inactivation of HBDH by MB. Fig. 5 shows that the response of these electrodes dropped by an unacceptable margin of approximately 7% after 14 weeks storage.

In summary, biosensor electrodes containing a mediator of the present invention displayed responses which were stable after at least 26 weeks storage. In contrast, those electrodes incorporating a traditional mediator such as MB which is an irreversible enzyme inhibitor exhibited responses which declined after only 14 weeks storage.

EXAMPLE 3

Evaluation of 1,10-PQ in Dry Strips containing glucose dehydrogenase (GDH):

Screen-printed electrodes were produced from an aqueous carbon ink 15 incorporating 1,10-PQ or MB at a level of 2.4 or 4.3 mg/g ink, respectively, together with the enzyme Glucose dehydrogenase (120 units/g ink) and NAD⁺ (110 mg/g ink). The ink also contained a polysaccharide binder.

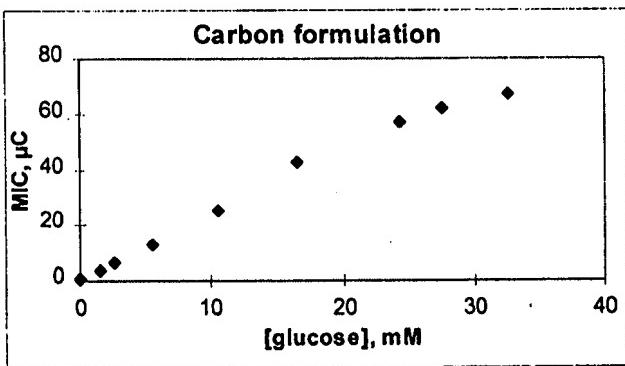
The calibrated dose response curve for the electrodes is given in Fig. 8. The electrodes were tested with whole blood containing physiologically relevant 20 concentrations of glucose ranging from 3.3 to 26 mM. A poise potential of +50 mV was maintained against a printed Ag/AgCl electrode. The electrodes produced a linear response over the glucose range. Thus, it was demonstrated that a mediator of the present invention can be used to construct a clinically useful glucose sensor which operates at a particularly low applied potential.

EXAMPLE 4

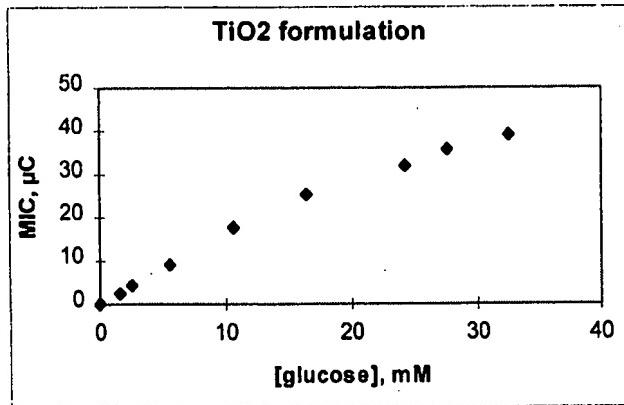
Electrode strips were prepared utilizing the construction illustrated in Figures 1 and 2 with a silver/silver chloride reference/counter electrode and a working electrode prepared by screen printing a formulation in accordance with Table 2. In one case, the filler was 25 weight percent ultra fine carbon and in the other case the

filler was 25 weight percent titania. In both cases the enzyme was Glucose Dehydrogenase (GDH), the cofactor was NAD, the mediator was 1, 10-PQ, the binder was guar gum, the protein stabilizer was Bovine serum albumin (BSA) and the buffer was Tris (.325 weight percent).

These electrode strips were evaluated by applying a 200 mV potential between the reference/counter electrode and the working electrode while an aqueous glucose solution covered both electrodes. The observed current from 15 to 20 seconds after the application of the potential was integrated and plotted against the glucose contents of the test solutions. The carbon-filled formulation gave a slope of 2.6 microcoulomb per mM of glucose and an X axis intercept of -1 microcoulomb while the titania-filled formulation gave a slope of 1.5 microcoulomb per mM of glucose and an X axis intercept of 0.6 microcoulomb. The plots were as follow:



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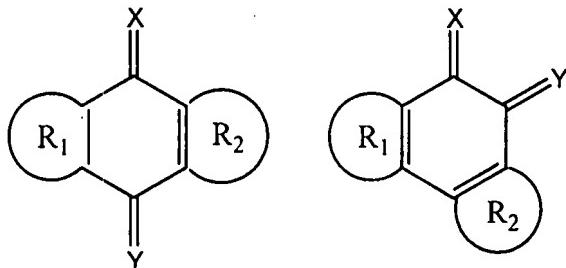


Other Embodiments

It is to be understood that while the invention has been described in conjunction with the detailed description thereof, that the foregoing description is intended to illustrate and not to limit the scope of the invention. Other aspects, 5 advantages, and modifications are within the scope of the invention.

Claims:

1. A single use disposable electrode strip for attachment to the signal readout circuitry of a sensor system to detect a current representative of an analyte in an aqueous sample, the strip comprising:
- 5 a) an elongated support having a substantially flat, planar surface, adapted for releasable attachment to said readout circuitry;
- 10 b) a first conductor extending along said surface and comprising a conductive element for connection to said readout circuitry;
- 15 an active electrode on said surface in contact with said first conductor, said active electrode comprising a nicotinamide cofactor-dependent enzyme, a nicotinamide cofactor, and a mediator compound having one of the following two formulae:



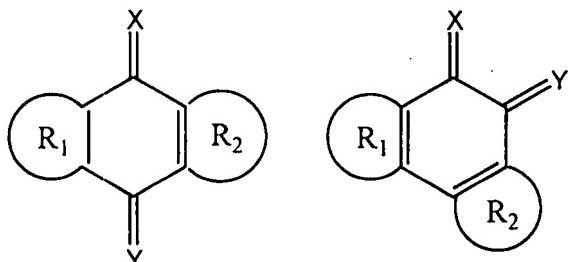
where X and Y can independently be oxygen, sulphur, CR^3R^4 , NR^3 , or NR^3R^{4+} or the functional group CZ^1Z^2 , where Z^1 and Z^2 are electron withdrawing groups; R_1 and R_2 can independently be a substituted or unsubstituted aromatic or heteroaromatic group; and R^3 and R^4 can independently be a hydrogen atom, a hydroxyl group or a substituted or unsubstituted alkyl, aryl, heteroaryl, amino, alkoxy, or aryloxy group,

20 wherein said active electrode is formulated with filler and binder ingredients such that said electrode gives a

monotonic response to concentrations of said analyte between about 1 and 8 mM when measurement is made in a kinetic mode in which simultaneous oxidation and reduction of the mediator occurs during the measurement;

- 5 c) a second conductor extending along said surface, comprising a conductive element for connection to said readout circuitry;
- d) a reference/counter electrode in contact with said second conductor;
- 10 e) said conductors being spaced apart so as not to be in electrical contact and being configured so as not to be brought into electrical contact when said aqueous sample is placed on said strip;
- f) said active electrode and said reference/counter electrode being configured so that both may be simultaneously covered by a small drop of said aqueous sample to provide an electrical conduction path between said electrodes.

2. A process of measuring the concentration in an aqueous sample of an analyte subject to oxidation by a NAD(P)⁺ dependent enzyme comprising
- 5 a) oxidizing the analyte with the NAD(P)⁺ dependent enzyme in the presence of NAD(P)⁺;
- oxidizing the NAD(P)H generated by reaction with the analyte and NAD(P)⁺ dependent enzyme with a mediator compound having one of the following two formulae:



10 where X and Y can independently be oxygen, sulphur, CR³R⁴, NR³, or NR³R⁴⁺ or the functional group CZ¹Z², where Z¹ and Z² are electron withdrawing groups; R₁ and R₂ can independently be a substituted or unsubstituted aromatic or heteroaromatic group; and R³ and R⁴ can independently be a hydrogen atom, a hydroxyl group or a substituted or unsubstituted alkyl, aryl, heteroaryl, amino, alkoxy, or aryloxy group; and

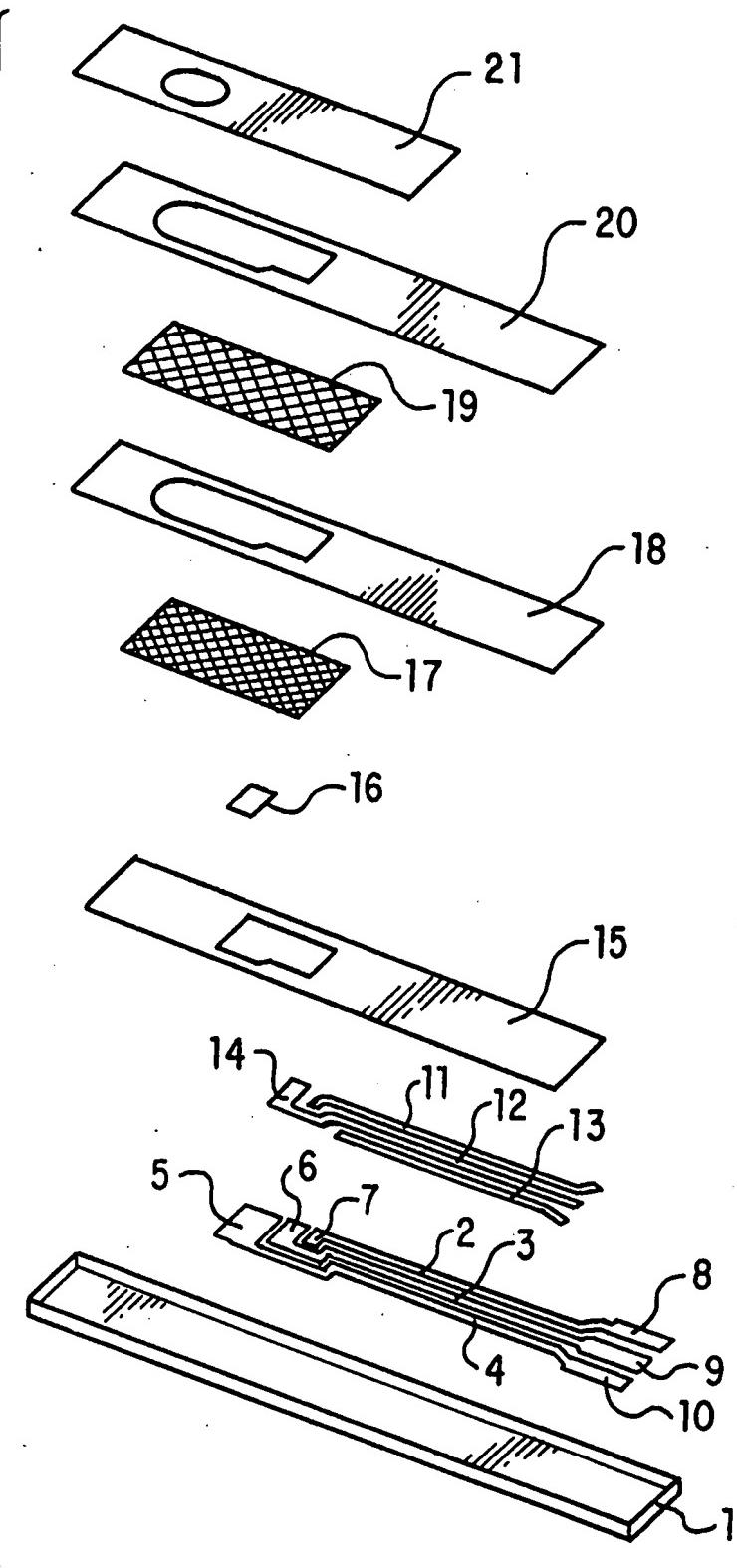
- 15 b) applying an electrical potential at an electrode to reoxidize the mediator compound reduced in oxidizing NAD(P)H and observing the resultant current,

wherein some of the mediator compound is being reduced by reaction with NAD(P)H while some of the mediator compound is being oxidized by transfer of electrons to said electrode during a measurement period and the rate of oxidation of the mediator compound over said measurement period and consequently the resultant observed current is monotonically related to the concentration of analyte in the sample.

3. The process of Claim 2 wherein the NAD(P)⁺ dependent enzyme, NAD(P)⁻ and mediator compound have been applied to the surface of said electrode in combination with a binder and a filler.
4. The process of Claim 3 wherein the current observed during the measurement period is linearly related to the concentration of the analyte in the sample.
5. The electrode strip of Claim 1 wherein the mediator compound is 1, 10-phenanthroline quinone.
6. The electrode strip of Claim 5 wherein the cofactor-dependent enzyme is Glucose Dehydrogenase.
- 10 7. The electrode strip of Claim 5 wherein the cofactor-dependent enzyme is 3-Hydroxybutyrate Dehydrogenase.
8. The process of Claim 2 wherein the mediator component is 1, 10-phenanthroline quinone.
9. The process of Claim 8 wherein the cofactor-dependent enzyme is Glucose Dehydrogenase.
- 15 10. The process of Claim 8 wherein the cofactor dependent enzyme is 3-Hydroxybutyrate Dehydrogenase.
11. The process of Claim 2 wherein the applied potential is 200 mV or less.

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FIG. 1



2/6

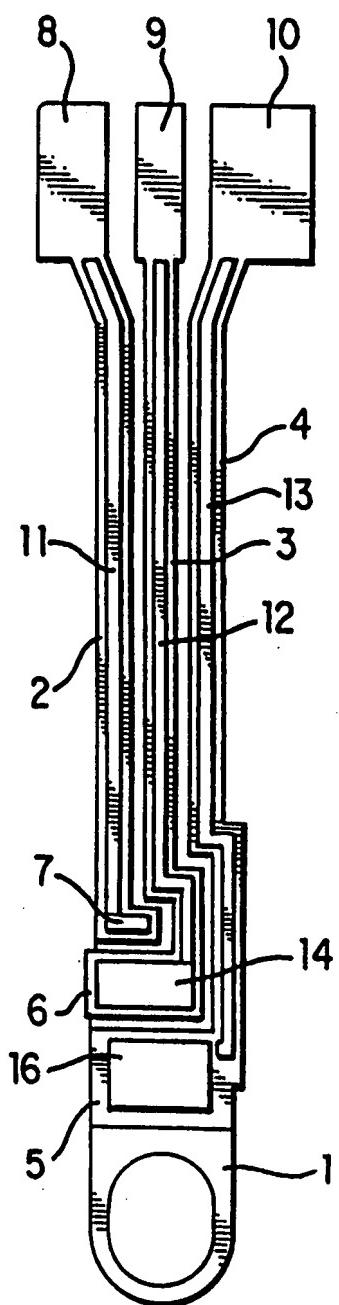


FIG. 2

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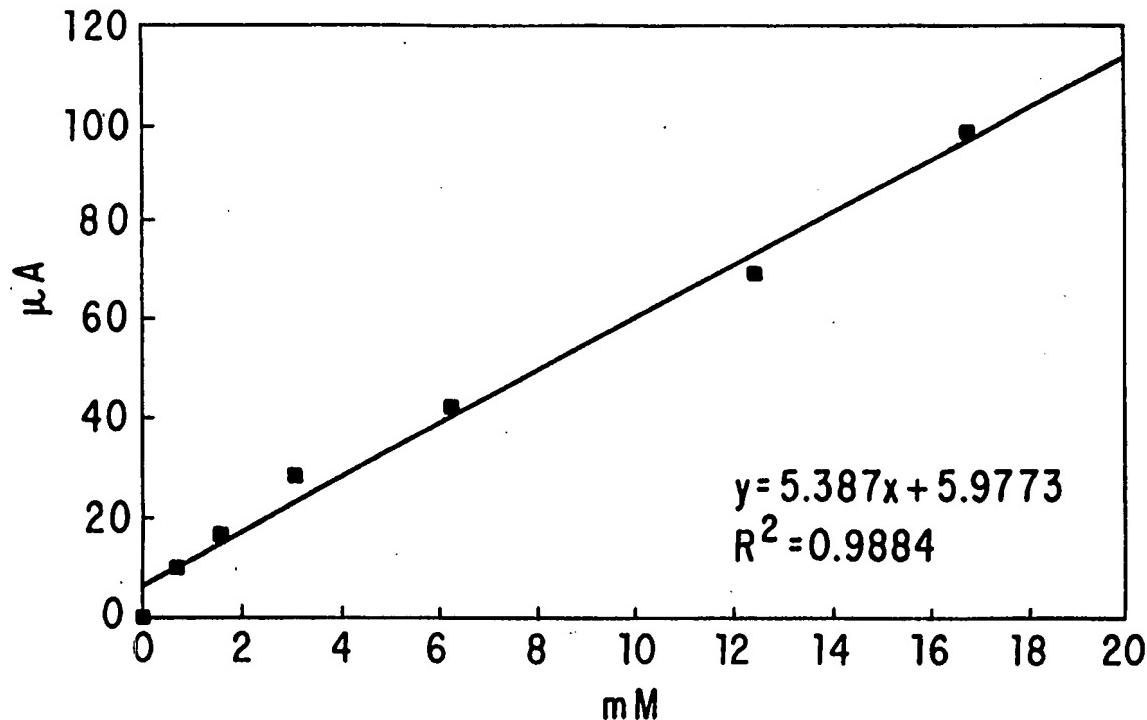


FIG. 3

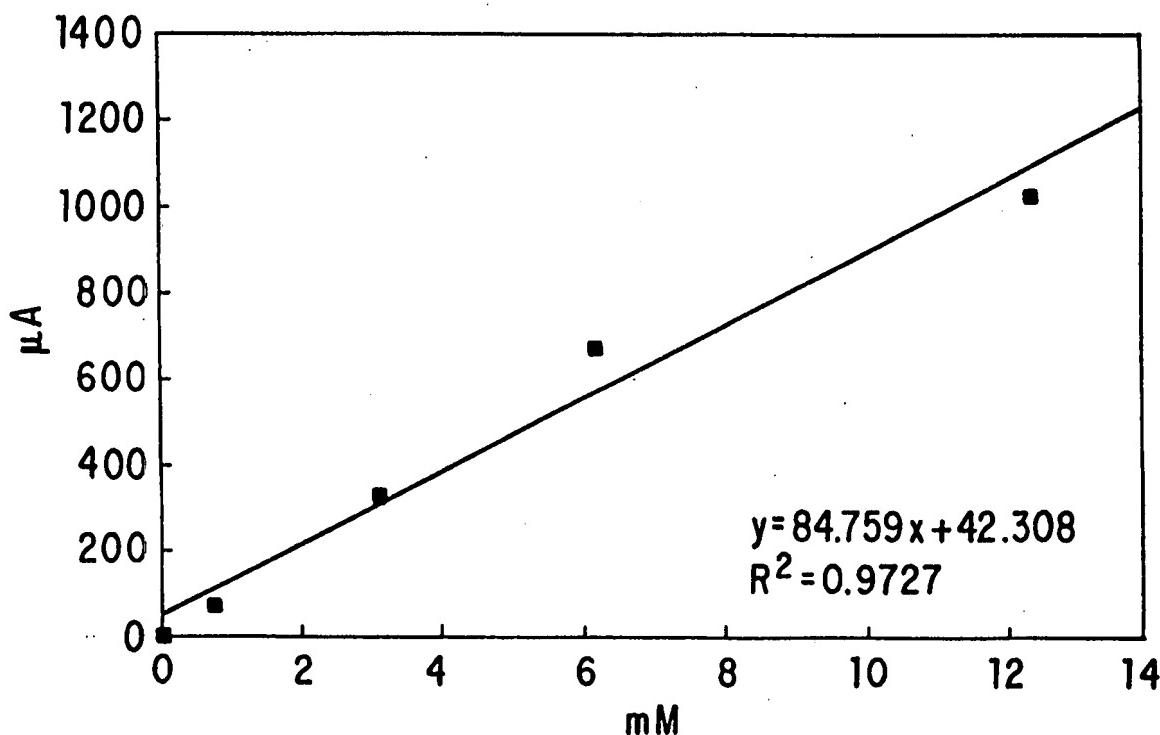


FIG. 4

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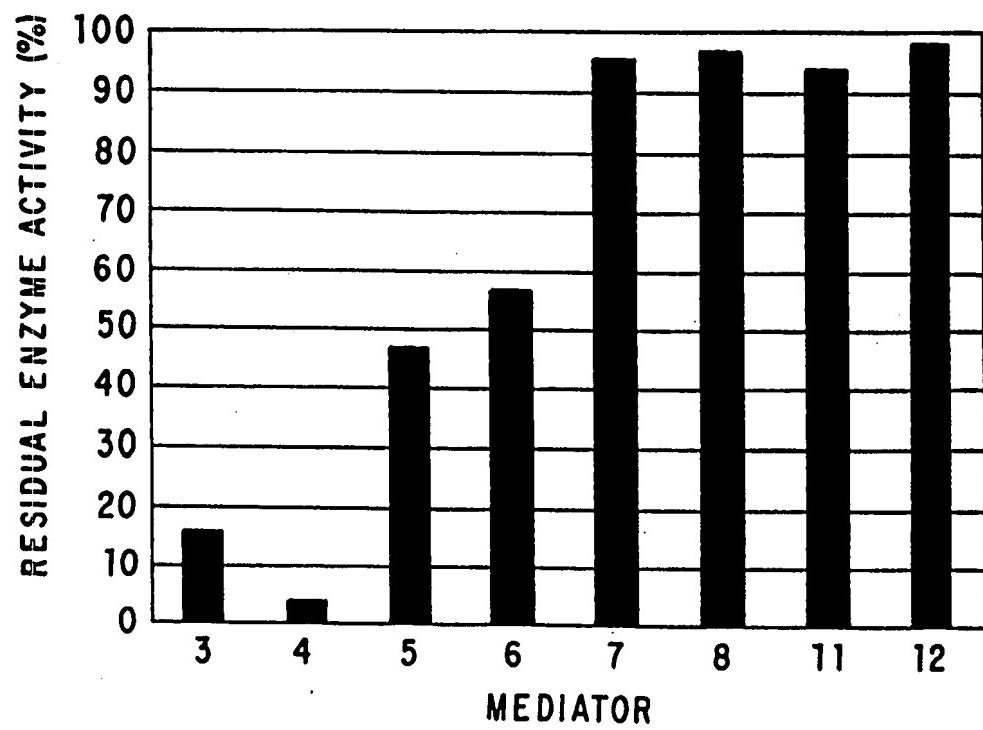


FIG. 5

5/6

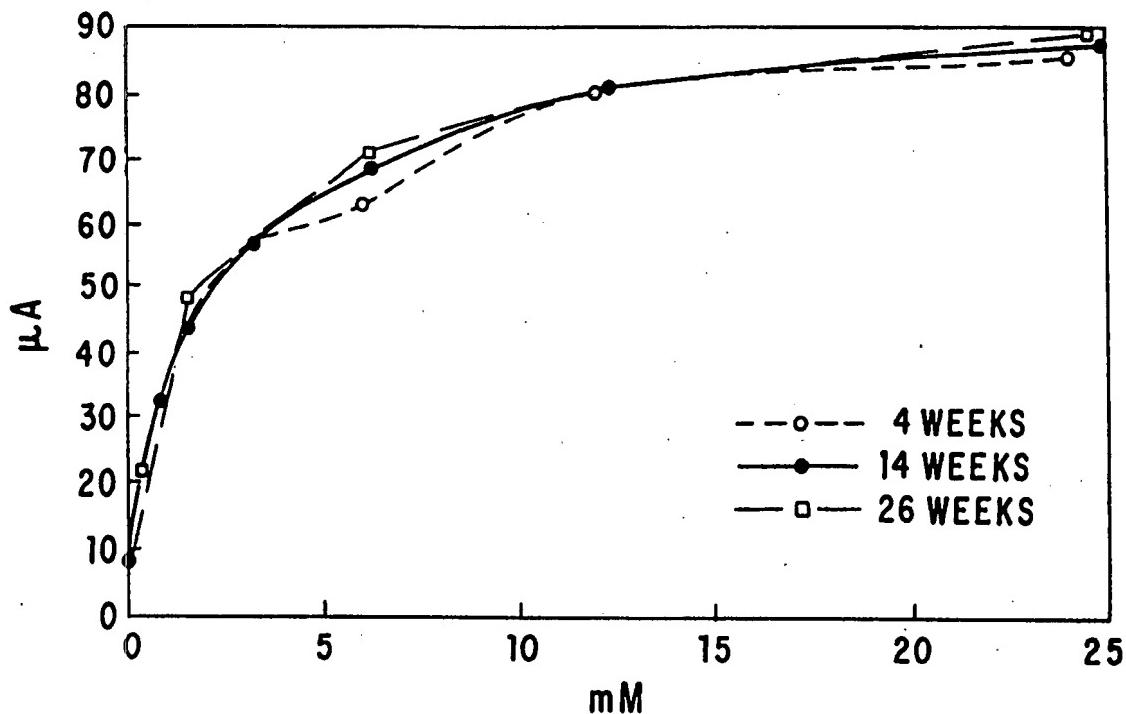


FIG. 6

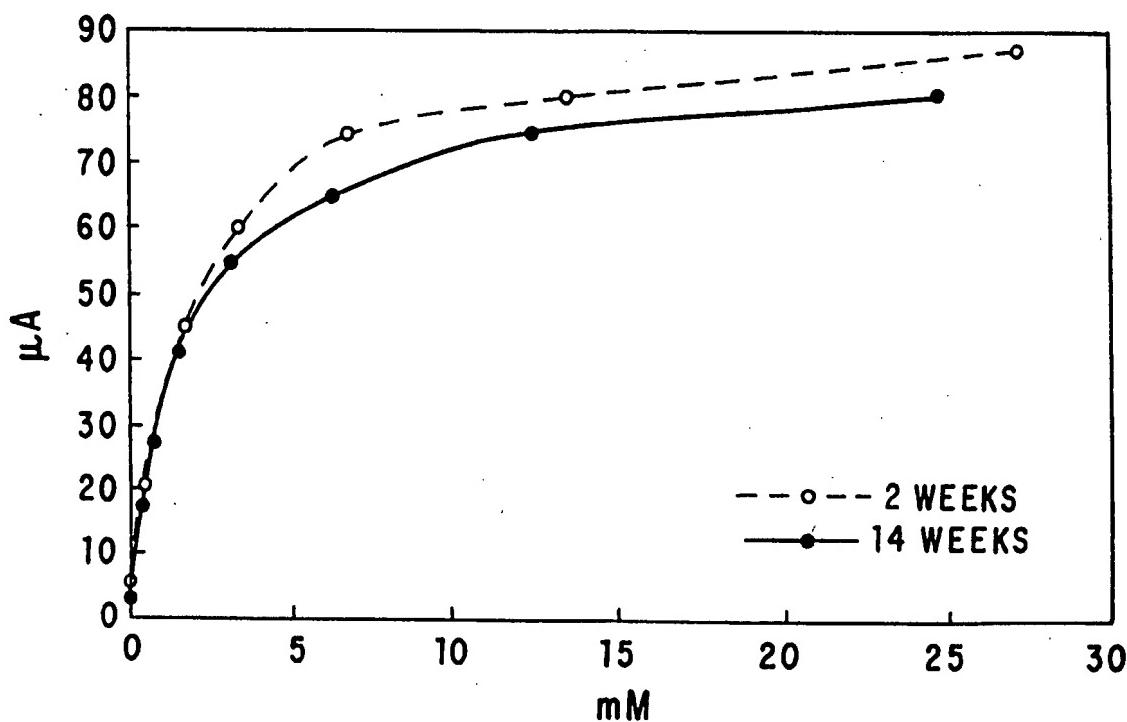


FIG. 7

6/6

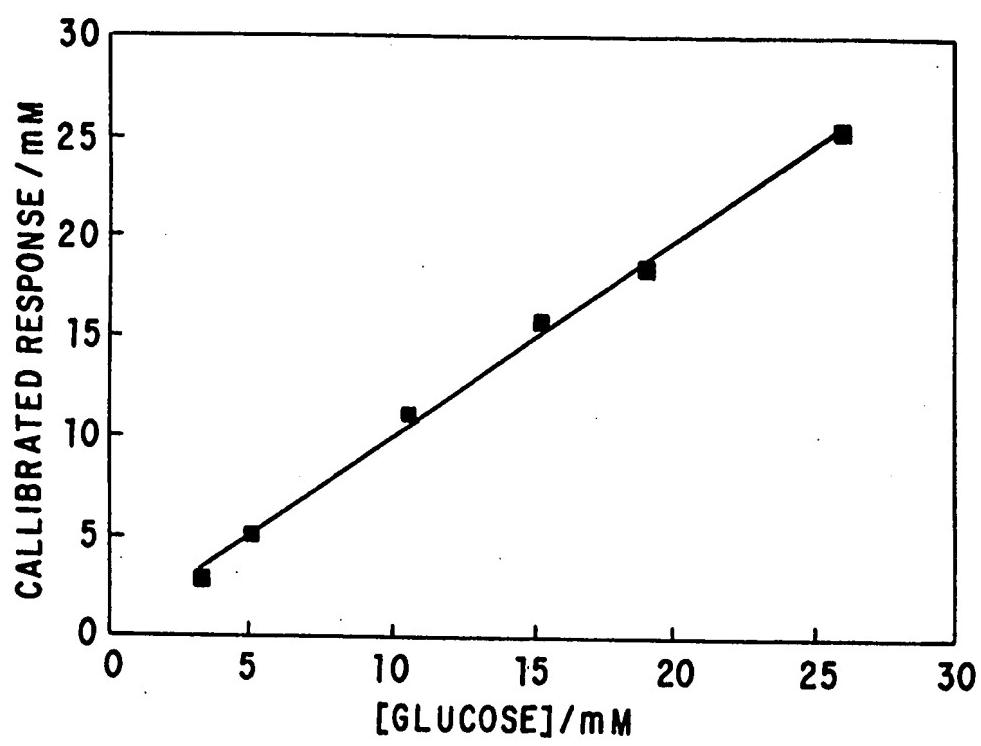


FIG. 8

INTERNATIONAL SEARCH REPORT

International Application No

PCT/US 98/21815

A. CLASSIFICATION OF SUBJECT MATTER

IPC 6 C12Q1/00 G01N27/30 G01N27/327

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC 6 G01N C12Q

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category °	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	GENG, L. ET AL.: "Amperometric biosensors based on dehydrogenase/NAD and heterocyclic quinones" BIOSENSORS & BIOELECTRONICS, vol. 11, no. 12, 1996, pages 1267-1275, XP002095074 cited in the application see the whole document ----	1-11
A	ECKERT, T. S. ET AL.: "Chemical Properties of Phenanthrolinequinones and the Mechanism of Amine Oxidation by o-Quinones of Medium Redox Potentials" JOURNAL OF THE AMERICAN CHEMICAL SOCIETY, vol. 105, 1983, pages 4431-4441, XP002095436 see the whole document ----	1-11
		-/-

Further documents are listed in the continuation of box C.

Patent family members are listed in annex.

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Date of the actual completion of the international search

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INTERNATIONAL SEARCH REPORT

International Application No

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Category	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
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A	US 4 758 323 A (DAVIS GRAHAM ET AL) 19 July 1988 see abstract; example 3 ---	1-4,6,7, 11
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A	US 5 628 890 A (CARTER NIGEL F ET AL) 13 May 1997 see figure 1 -----	1,2

INTERNATIONAL SEARCH REPORT

Information on patent family members

International Application No

PCT/US 98/21815

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			DE 3486221 T		27-01-1994
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FOR

IMPROVED ELECTROCHEMICAL CELL

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Publication date: 2001-10-16

Inventor:

Applicant:

Classification:

- International: **G01N27/30; G01N27/327; G01N27/403; G01N27/416; G01N27/30; G01N27/327; G01N27/403; G01N27/416;**
(IPC1-7): G01N27/30; G01N27/416

- European: G01N27/30F; G01N27/403

Application number: JP19980543209T 19980325

Priority number(s): AU1997PO05857 19970325; WO1998AU00200
19980325

Also published as:



WO9843074 (A1)



EP1012590 (A1)



US6454921 (B1)



EP1012590 (A4)



EP1012590 (A0)

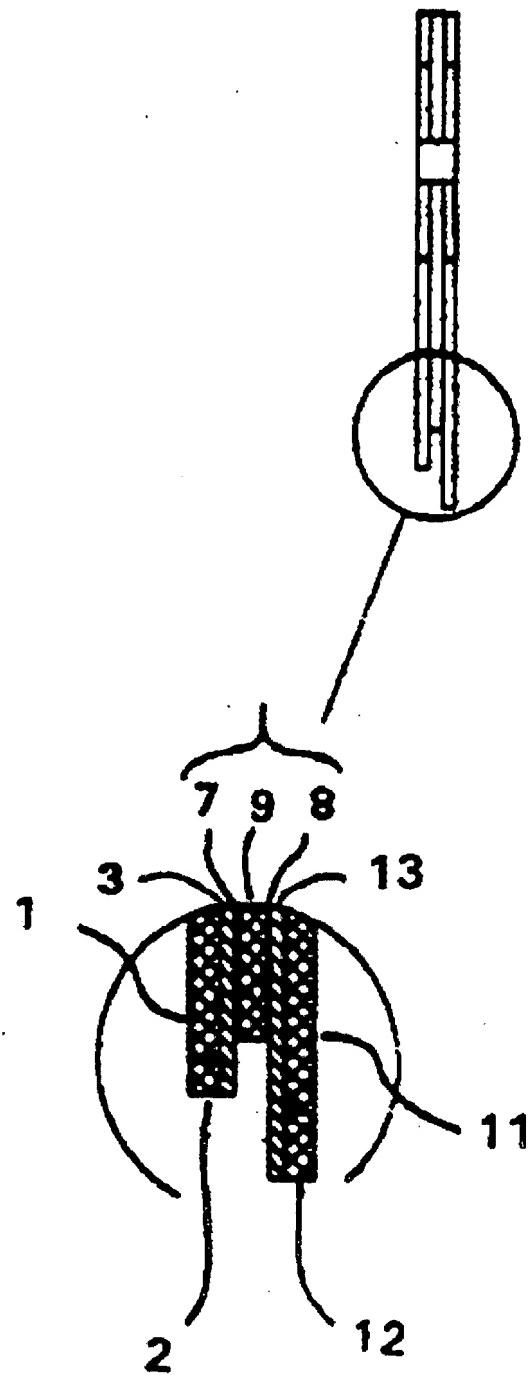
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Abstract not available for JP2001519032T

Abstract of corresponding document: **WO9843074**

The invention relates to an amperometric electrochemical cell having a first insulating substrate (1) carrying a first electrode (2), a second insulating substrate (11) carrying a second electrode (10), said electrodes being disposed to face each other and spaced apart by less than 500 μm , and defining a sample reservoir (4) therebetween, and wherein at least one, and preferably both, insulating substrates and the electrode carried thereon include an electromagnetic radiation transmissive portion in registration with said reservoir. The walls of the electrochemical cell (2, 12) may be formed from a thin metallic portion (3) on a transparent substrate (1). Such cells are useful in providing visual confirmation of the validity of the electrochemical measurement.



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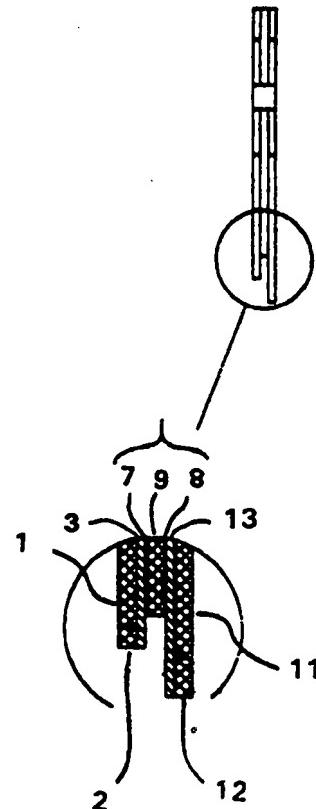
INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(51) International Patent Classification ⁶ : G01N 27/30	A1	(11) International Publication Number: WO 98/43074 (43) International Publication Date: 1 October 1998 (01.10.98)
(21) International Application Number: PCT/AU98/00200		(81) Designated States: AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ, DE, DK, EE, ES, FI, GB, GE, GH, GM, GW, HU, ID, IL, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, UA, UG, US, UZ, VN, YU, ZW, ARIPO patent (GH, GM, KE, LS, MW, SD, SZ, UG, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, ML, MR, NE, SN, TD, TG).
(22) International Filing Date: 25 March 1998 (25.03.98)		
(30) Priority Data: PO 5857 25 March 1997 (25.03.97) AU		
(71) Applicant (<i>for all designated States except US</i>): MEMTEC AMERICA CORPORATION [US/US]; Suite 700, 9690 Deereco Road, Timonium, MD 21093 (US).		
(72) Inventors; and		Published
(75) Inventors/Applicants (<i>for US only</i>): HODGES, Alastair, McIndoe [AU/AU]; 15 Jasmine Court, Blackburn South, VIC 3130 (AU). BECK, Thomas, William [AU/AU]; 31 Drummond Street, South Windsor, NSW 2756 (AU). MAXWELL, Ian, Andrew [AU/AU]; 3 Whiting Street, Leichhardt, NSW 2040 (AU).		<i>With international search report.</i>
(74) Agent: SHELSTON WATERS; 60 Margaret Street, Sydney, NSW 2000 (AU).		

(54) Title: IMPROVED ELECTROCHEMICAL CELL

(57) Abstract

The invention relates to an amperometric electrochemical cell having a first insulating substrate (1) carrying a first electrode (2), a second insulating substrate (11) carrying a second electrode (10), said electrodes being disposed to face each other and spaced apart by less than 500 µm, and defining a sample reservoir (4) therebetween, and wherein at least one, and preferably both, insulating substrates and the electrode carried thereon include an electromagnetic radiation transmissive portion in registration with said reservoir. The walls of the electrochemical cell (2, 12) may be formed from a thin metallic portion (3) on a transparent substrate (1). Such cells are useful in providing visual confirmation of the validity of the electrochemical measurement.



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"IMPROVED ELECTROCHEMICAL CELL"

Technical Field

This invention relates to disposable electrochemical sensors of the type used for
5 quantitative analysis, for example, of glucose levels in blood, or the like.

Background Art

Light transmissive electrodes are known in the prior art, however they have not previously been applied to amperometric cells. For example, GB 2 194 112 discloses the use of optically transparent electrodes used to drive a microelectrophoresis cell while laser
10 doppler velocimetry is used to determine the velocity and micro current motion of charged particles within the sample.

JP, A 05080018 discloses another approach to making transparent electrodes by the use of conductive glass for electrochromic and field emission devices.

JP, A 06310746 also teaches the use and formation of yet another type of
15 conducting transparent electrode formed from the deposition of organic conducting polymers onto a glass slide. This type of electrode is useful in solar energy collection cells. Ullery, in US 4,254,546 also discloses a photovoltaic cell in which the top layer is a light collecting electrode.

US 4,782,265 discloses two spaced apart translucent electrodes useful in luminescent
20 cells. However, US 4,782,265 specifically teaches that gold, silver, aluminium, platinum and the like are only suitable for the production of non-transmissive electrodes.

In co-pending applications PCT/AU96/00207, PCT/AU96/00365, PCT/AU96/00723 and PCT/AU96/00724 (the contents of which are incorporated herein by reference) there are described various very thin electrochemical sensors or cells. These cells are by a pair

of oppositely facing spaced apart electrodes which are formed as thin metal coatings (for example sputter coatings) deposited on thin inert plastic film (for example 100 micron thick PET). The electrodes are separated one from the other by a spacer of thickness of for example 500 μm or less.

5 Such cells may be provided with one or more fluid passageways into and out of the sample reservoir whereby the cell may be filled with an analyte and air expelled during filling. In some embodiments the analyte is drawn into the cell by the energy liberated as a reagent contained therein dissolves.

The sensors are, as discussed above, very small and normally contain only small
10 amounts of the liquid sample. Accurate measurement requires that the cell be filled with liquid. Even minute variations in the quantity of liquid in the cell can effect the sensing measurements. It can be difficult for a user to be sure that in use the cell has been uniformly filled with a sample to be analysed.

Further, sensors of the kind under discussion are usually intended to be discarded
15 after use. If a user is distracted after use or prior to disposal it is not always easy for the user to know which sensors have been used and which have not.

It is an object of the present invention to overcome or ameliorate at least one of the disadvantages of the prior art, or to provide a useful alternative.

Disclosure of the Invention

20 According to one aspect the present invention provides an amperometric electrochemical cell comprising a first insulating substrate carrying a first electrode, a second insulating substrate carrying a second electrode, said electrodes being disposed to face each other and spaced apart by less than 500 μm , and defining a sample reservoir therebetween, wherein at least one of said insulating substrates and the electrode carried

thereon includes an electromagnetic radiation transmissive portion in registration with said reservoir.

Preferably, both said insulating substrates and the electrodes thereon include a transmissive portion. Most preferably, the transmissive portion is formed by a conductive 5 metallic coating on the substrate, which is of a thickness such that it is transparent or translucent. Suitable substances for the metallic coating include gold, indium oxide, tin oxide or mixtures thereof. A suitable substrate is PET.

According to a second aspect, the invention provides a method of filling an amperometric cell comprising the steps of:

- 10 a) drawing a liquid sample into said cell comprising a first insulating substrate carrying a first electrode, a second insulating substrate carrying a second electrode, said electrodes being disposed to face each other and spaced apart by less than 500 µm, and defining a sample reservoir therebetween, wherein at least one of said insulating substrates and the electrode carried thereon includes an electromagnetic radiation transmissive portion in
15 registration with said reservoir
- b) exposing the transmissive portion to electromagnetic radiation
- c) monitoring a property of the electromagnetic radiation passing and/or reflected through
 said transmissive portion
- d) comparing said monitored property with a predetermined value indicative of the cell
20 being filled, and
- e) continuing to draw the liquid sample into the cell until said monitored property reaches
 said predetermined value.

According to a third aspect, the invention provides a method of determining whether an amperometric cell is filled with a liquid sample, said cell comprising a first insulating

- 4 -

- substrate carrying a first electrode, a second insulating substrate carrying a second electrode, said electrodes being disposed to face each other and spaced apart by less than 500 µm, and defining a sample reservoir therebetween, wherein at least one of said insulating substrates and the electrode carried thereon includes an electromagnetic radiation transmissive portion in registration with said reservoir, said method comprising
- 5 the steps of:
- a) exposing said transmissive portion to electromagnetic radiation,
 - b) monitoring a property of the electromagnetic radiation passing and/or reflected through said transmissive portion, and
- 10 c) comparing said monitored property with a predetermined value indicative of the cell being filled.

Suitable forms of electromagnetic radiation include visible, ultraviolet, infra-red and laser light. Daylight is especially preferred. The monitored property can include optical density, wavelength, refractive index and optical rotation.

- 15 In preferred embodiments, it is envisaged that the sample will be blood. The electromagnetic property may be monitored inside (for instance with a fibre optical device) or outside the cell, and the electromagnetic radiation may pass substantially directly through the cell or be internally reflected within the cell.

- According to a fourth aspect, the invention consist in a method for monitoring an
- 20 analyte in a liquid sample comprising the steps of :

- a) drawing the sample into an amperometric electrochemical cell comprising a first insulating substrate carrying a first electrode, a second insulating substrate carrying a second electrode, said electrodes being disposed to face each other and spaced apart by less than 500 µm, and defining a sample reservoir therebetween, wherein at least one of said

- 5 -

insulating substrates and the electrode carried thereon includes an electromagnetic radiation transmissive portion in registration with said reservoir,

- b) exposing the transmissive portion to electromagnetic radiation
- c) monitoring a property of the electromagnetic radiation passing and/or reflected
- 5 through said transmissive portion
- d) comparing said monitored property with a predetermined value indicative of the cell being filled, and
- e) prior to, simultaneously with or after any one of steps b) to d) applying a potential across the electrochemical cell and measuring the resultant current to detect the
- 10 analyte.

The method of the above aspect may also further comprise the step of:

- f) repeating steps a) to e) until the monitored property reaches the predetermined value.

In one preferred embodiment, the method is repeated on different cells, with blood as

15 the sample and visible light the electromagnetic radiation, until a valid measurement is obtained for blood glucose.

In a fifth aspect, the invention provides an apparatus for determining whether an amperometric cell according to the first aspect is filled with a liquid sample, said apparatus comprising an electromagnetic radiation means adapted to expose said transmissive portion

20 of said cell to electromagnetic radiation,

a monitoring means adapted to monitor a property of the electromagnetic radiation passing and/or reflected through said transmissive portion, and

a means for determining whether said monitored property has reached a predetermined value indicative of the cell being filled.

Preferably, said apparatus may also include means to apply potential across the amperometric cell and detect the resultant current. It may also include a validation means to confirm the cell is filled with a liquid sample.

- "Comprising" as herein used is used in an inclusive sense, that is to say in the sense
5 of "including" or "containing". The term is not intended in an exclusive sense
("consisting" of or "composed of").

Light-transmissive cells intended for spectrophotometric use are well known. However this has not previously been accomplished in a cell wherein the only surfaces suitable for a window are entirely covered by a metal electrode. One skilled in the art will
10 appreciate that whilst the embodiments of the invention are described with respect to light transmissive conductive coatings, such coatings may be also be transparent to some other forms of electromagnetic radiation which are not visible to the human eye.

Best Mode for Carrying out the Invention

The invention will now be particularly described by way of example only reference
15 to the accompanying schematic drawings wherein:

Figure 1 shows a cross section of a wall of a cell according to the present invention.

Figure 2 shows a cell according to one embodiment of the present invention.

Figure 3 shows a cross section of the cell in Figure 2.

An embodiment of the invention will now be described by way of example only.
20 Referring firstly to figure 1, each wall of the cell 2,12 comprises an insulating substrate 1,11 carrying an electrode 3,13 thereon.

The embodiment is generally in accordance with the apparatus described in our co-pending application PCT/AU96/00724 which is incorporated herein by reference. The apparatus hereof corresponds substantially to the apparatus described in that application

with the difference that electrode layer 3,13 which in application PCT/AU96/00724 was sputter coated palladium having a thickness of 100-1000 Angstrom, is replaced according to the present invention, by a light-transmissive conductive metallic coating of a thickness such that it is transparent or translucent. Gold, indium oxide, tin oxide and mixtures of 5 indium and tin oxides or other suitable light-transmissive conductive metallic coating may be utilised. Those skilled in the art will appreciate that transparent inorganic and organic polymers or mixtures thereof could also be used. Substrate 1 is also light-transmissive.

In one embodiment, the cell takes the form as shown in Figures 2 or 3. The cell comprises a first insulating substrate 1 consisting of a Melinex® PET layer, a first electrode 10 3 consisting of a conductive metallic layer on substrate 1, an adhesive layer 7, a PET spacer 9, a second adhesive layer 8, a second electrode 13 formed as a metallic layer formed as a metallic coating on second insulating substrate 11. Spacer 9 defines a sample reservoir 4 having a thickness corresponding to the thickness of the spacer 9 together with the thickness of adhesive layers 7 and 8. Access to the sample reservoir 4 is provided at 15 the side edge of the cell by notches 6.

At least one of the said insulating substrates and the electrode carried thereon includes an electromagnetic radiation transmissive portion 20 in registration with the reservoir 4.

In preferred embodiments of the invention, a sample to be analysed is introduced 20 to the cell by capillary action. The sample is placed on contact with notch 6 and is spontaneously drawn by capillary action into the reservoir 4, displaced air from the reservoir 4 venting from the opposite notch 6. A surfactant may be included in the capillary space to assist in drawing in the sample.

The cell is provided with connection means for example edge connectors whereby the cell may be placed into a measuring circuit. In a preferred embodiment this is achieved by making spacer 9 shorter than cell walls 2, 12 and by making one wall 2 of shorter length than the other 12. This forms a socket region having contact areas electrically connected

5 with the working and counter electrodes of a sensing apparatus. A simple tongue plug having corresponding engaging conduct surfaces can then be used for electrical connection.

Connectors of other form may be devised.

Chemicals for use in the cell may be supported on the cell electrodes or walls, may be supported on an independent support contained within the cell or may be self-

10 supporting.

In use, when the cell is filled with the liquid sample, e.g. blood, a film of the sample covers the inside of the transmissive portion 20 formed by substrate 1,11 and metal electrode 3,13 over reservoir 4, thereby indicating to the user when the cell is adequately filled, and clearly differentiating a used sensor from an unused one.

15 Apart from simple visual inspection, a user can also monitor the filling of the cell by exposing the electromagnetic radiation transmissive portion 20 to electromagnetic radiation such as infra-red, ultraviolet light or laser light and monitoring a property of the electromagnetic radiation (for example, optical density, colour or optical rotation) as it exits the cell, either by another transmissive portion on the opposite side of the reservoir, or

20 as a result of internal reflection within the reservoir.

A particular embodiment of an apparatus suitable for carrying out the inventive method is hereby described. The apparatus has means for holding and orienting a cell according to the present invention and exposing the electromagnetic portion of the cell to allow an electromagnetic radiation source, e.g. a light beam, to enter the reservoir.

Optionally, the apparatus may also be equipped with means for automatically placing the cells into the said holding means.

The path of the light beam is substantially linear, and it exits the cell either via another transparent portion of the cell, or is reflected back through the first mentioned 5 transmissive portion. The electromagnetic radiation leaving the cell is monitored by an appropriate detector. This detector monitors a property of the electromagnetic radiation leaving the cell for comparison with a predetermined value which is indicative of the cell being filled, for example if the cell is filled with blood up to or above the transmissive portion, a reduction in optical density will be detected. If the cell is empty or only partially 10 filled to this point, the optical density will remain high but will reduce as the reservoir is filled until it reaches the predetermined value indicating a full cell. It will be appreciated that the apparatus could be made to continue filling until the cell was satisfactorily filled, by means of a feedback system. The apparatus could also be adapted so that it performed the necessary electrochemical measurements on the cell, thus reducing the need for 15 excessive sample movement.

The apparatus preferably also includes a validation means which is triggered when it is determined that the cell is full and the sensing measurement can be accepted as valid.

As will be apparent to those skilled in the art from the teaching hereof the invention may be embodied in other forms without departing herefrom.

CLAIMS

1. An amperometric electrochemical cell comprising a first insulating substrate carrying a first electrode, a second insulating substrate carrying a second electrode, said electrodes being disposed to face each other and spaced apart by less than 500 µm, and defining a sample reservoir therebetween, wherein at least one of said insulating substrates and the electrode carried thereon includes an electromagnetic radiation transmissive portion in registration with said reservoir.
2. An amperometric electrochemical cell according to claim 1 wherein both of said insulating substrates and the electrodes carried thereon include a transmissive portion.
- 10 3. An amperometric electrochemical cell according to claim 1 or claim 2 wherein the electromagnetic radiation transmissive portion of each electrode is formed by a conductive metallic coating on said substrate(s).
4. An amperometric electrochemical cell according to claim 3 wherein the conductive metallic coating is of a thickness such that it is transparent or translucent.
- 15 5. An amperometric electrochemical cell according to claims 3 or 4 wherein the metallic coating is selected from one or more members of the group consisting of: gold, indium oxide, tin oxide and mixtures thereof.
6. An amperometric electrochemical cell according to any one of the preceding claims wherein the insulating substrate(s) is/are PET.
- 20 7. A method of filling an amperometric cell comprising the steps of:
 - a) drawing a liquid sample into said cell comprising a first insulating substrate carrying a first electrode, a second insulating substrate carrying a second electrode, said electrodes being disposed to face each other and spaced apart by less than 500 µm, and defining a sample reservoir therebetween, wherein at least one of said insulating substrates and the

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electrode carried thereon includes an electromagnetic radiation transmissive portion in registration with said reservoir

b) exposing the transmissive portion to electromagnetic radiation

c) monitoring a property of the electromagnetic radiation passing and/or reflected through

5 said transmissive portion

d) comparing said monitored property with a predetermined value indicative of the cell being filled, and

e) continuing to draw the liquid sample into the cell until said monitored property reaches said predetermined value.

10 8. A method of determining whether an amperometric cell is filled with a liquid sample, said cell comprising a first insulating substrate carrying a first electrode, a second insulating substrate carrying a second electrode, said electrodes being disposed to face each other and spaced apart by less than 500 µm, and defining a sample reservoir therebetween, wherein at least one of said insulating substrates and the electrode carried thereon includes
15 an electromagnetic radiation transmissive portion in registration with said reservoir, said method comprising the steps of:

a) exposing said transmissive portion to electromagnetic radiation,

b) monitoring a property of the electromagnetic radiation passing and/or reflected through said transmissive portion, and

20 c) comparing said monitored property with a predetermined value indicative of the cell being filled.

9. A method according to claim 7 or 8 wherein the electromagnetic radiation is selected from the group consisting of visible light, ultraviolet, infra-red, laser

10. A method according to claim 9 wherein the visible light is daylight

11. A method according to any one of claims 7 to 10 wherein said monitored property is selected from the group consisting of optical density, wavelength, refractive index and optical rotation.
12. A method according to any one of claims 7 to 11 wherein the liquid sample is
5 blood.
13. A method according to any one of claims 7 to 12 wherein the electromagnetic property is monitored outside the cell.
14. A method according to any one of claims 7 to 12 wherein the electromagnetic property is monitored inside the cell.
- 10 15. A method according to claim 13 wherein the electromagnetic radiation passing substantially directly through the sample is monitored.
16. A method according to claim 13 wherein the electromagnetic radiation reflected internally in the cell is monitored.
17. A method according to claim 14 wherein the electromagnetic property is observed
15 by a fibre optical device.
18. A method for monitoring an analyte in a liquid sample comprising the steps of :
 - a) drawing the sample into an amperometric electrochemical cell comprising a first insulating substrate carrying a first electrode, a second insulating substrate carrying a second electrode, said electrodes being disposed to face each other and spaced apart by less than 500 µm, and defining a sample reservoir therebetween, wherein at least one of said insulating substrates and the electrode carried thereon includes an electromagnetic radiation transmissive portion in registration with said reservoir,
 - b) exposing the transmissive portion to electromagnetic radiation

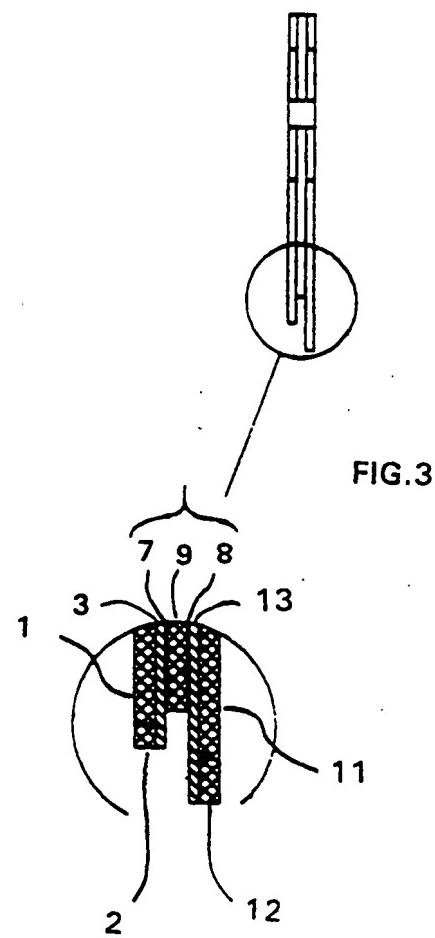
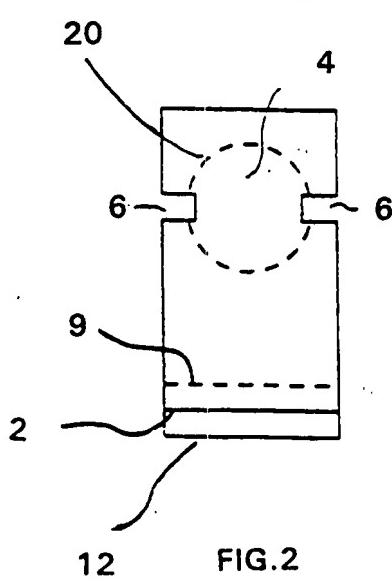
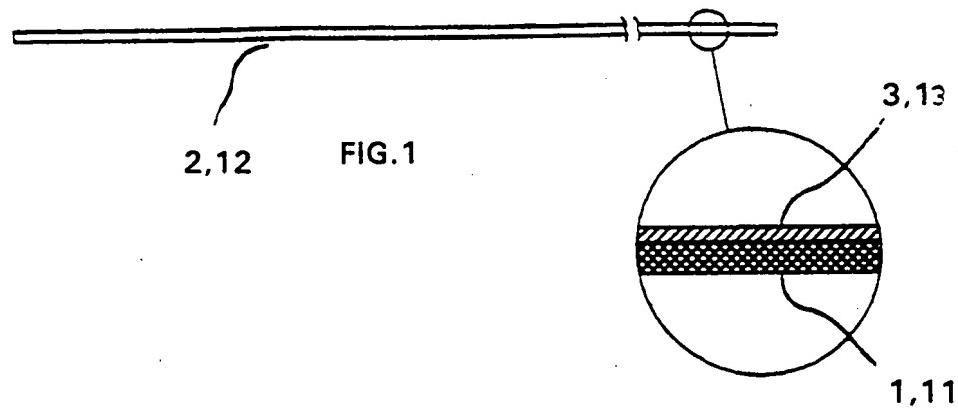
- c) monitoring a property of the electromagnetic radiation passing and/or reflected through said transmissive portion
 - d) comparing said monitored property with a predetermined value indicative of the cell being filled, and
- 5 e) prior to, simultaneously with or after any one of steps b) to d) applying a potential across the electrochemical cell and measuring the resultant current to detect the analyte.
19. A method according to claim 18 further comprising the step of
- f) repeating steps a) to e) until the monitored property reaches the predetermined
- 10 value.
20. A method according to claim 19 wherein steps a) to e) are repeated on the same cell.
21. A method according to claim 19 wherein steps a) to e) are repeated on a new cell.
22. A method according to claim 19 wherein the sample is blood.
- 15 23. A method according to claim any one of claims 18 to 22 wherein the electromagnetic radiation is visible light.
24. A method according to any one of claims 18 to 23 wherein the monitored property is selected from the group consisting of optical density, wavelength, refractive index and optical rotation.
- 20 25. A method according to any one of claims 18 to 24 wherein the analyte is glucose.
26. An apparatus for determining whether an amperometric cell according to any one of claims 1 to 6 is filled with a liquid sample, said apparatus comprising an electromagnetic radiation means adapted to expose said transmissive portion of said cell to electromagnetic radiation,

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a monitoring means adapted to monitor a property of the electromagnetic radiation passing and/or reflected through said transmissive portion, and
a means for determining whether said monitored property has reached a predetermined value indicative of the cell being filled.

- 5 27. An apparatus according to claim 26 further including means to apply a potential across the amperometric cell and detect the resultant current.
28. An apparatus according to claims 26 or 27 further including a validation means to confirm that the cell is filled with liquid sample.

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INTERNATIONAL SEARCH REPORT

International Application No.

PCT/AU 98/00200

A. CLASSIFICATION OF SUBJECT MATTER

Int Cl⁶: G01N 27/30

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC G01N 27/26, 27/28, 27/30, 27/403, 27/404

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

AU : IPC as above

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)

DERWENT : (TRANSPAREN: OR TRANSLUC: OR GLASS# OR PLASTIC OR CLEAR) AND
(SUBSTRATE# OR ELECTRODE#)

JAPIO : as per DERWENT

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	GB 2194112 A (MALVERN INSTRUMENTS LIMITED) 24 February 1988 abstract, page 1 lines 76-82, page 2 lines 6-14, Fig 1	1-28
X	Derwent Abstract Accession No. 93-140898/17, class S03, JP,A, 05080018 (RIKAGAKU KENKYUSHO) 30 March 1993 abstract	1-28

Further documents are listed in the continuation of Box C

See patent family annex

* Special categories of cited documents:

- "A" document defining the general state of the art which is not considered to be of particular relevance
- "E" earlier document but published on or after the international filing date
- "L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)
- "O" document referring to an oral disclosure, use, exhibition or other means
- "P" document published prior to the international filing date but later than the priority date claimed

"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone

"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art

"&" document member of the same patent family

Date of the actual completion of the international search

26 May 1998

Date of mailing of the international search report

-2 JUN 1998

Name and mailing address of the ISA/AU
AUSTRALIAN PATENT OFFICE
PO BOX 200
WODEN ACT 2606
AUSTRALIA
Facsimile No.: (02) 6285 3929

Authorized officer

SOOSA GNANASINGHAM

Telephone No.: (02) 6283 2172

INTERNATIONAL SEARCH REPORT

International Application No.

PCT/AU 98/00200

DOCUMENTS CONSIDERED TO BE RELEVANT		
Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	Derwent Abstract Accession No. 95-026336/04, class S03, JP,A, 06310746 (HITACHI LTD) 4 November 1944 abstract	1-28
X	US 4782265 A (SCHAPER et al) 1 november 1988 column 1 lines 6-13, column 2 lines 14-34, claim 1	1-28
X	US 4254546 A (ULLERY, Jr) 10 March 1981 column 3 lines 11-15, column 9 lines 49-52 (claim 1)	1-28
X,P	Patent Abstracts of Japan, JP, 9-243588 A (MATSUSHITA ELECTRIC IND CO LTD) 19 September 1997 abstract	1-28
X,P	Patent Abstracts of Japan, JP, 9-222408 A (MATSUSHITA ELECTRIC IND CO LTD) 26 August 1997 abstract	1-28
X,P	Patent Abstracts of Japan, JP, 9-236570 A (MATSUSHITA ELECTRIC IND CO LTD) 9 September 1997 abstract	1-28
X,P	DE 29709141 U1 (SCHWABE-INST MESS & SENSOR TECHNIK EV) 28 August 1997 whole document	1-28
Y	EP 585933 A2 (MATSUSHITA ELECTRIC INDUSTRIAL CO LTD) 9 March 1994 column 6 lines 28-40, claim 4	1-28

INTERNATIONAL SEARCH REPORT

Information on patent family members

International Application No.

PCT/AU 98/00200

This Annex lists the known "A" publication level patent family members relating to the patent documents cited in the above-mentioned international search report. The Australian Patent Office is in no way liable for these particulars which are merely given for the purpose of information.

Patent Document Cited in Search Report			Patent Family Member				
GB	2194112	DE	3723388	JP	63045553		
US	4254546	BE	861439	BR	7707957	CA	1089946
		DE	2751393	ES	464679	FR	2373165
		GB	1559247	IL	53276	IN	148687
		IT	1091265	JP	53070781	MX	144753
		NL	7712838	ZA	7706512	US	4127424

END OF ANNEX